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(54) Novel peptidase inhibitors.

(57) This invention relates to analogs of peptidase substrates in which the amide group containing the scissile amide bond of the substrate peptide has been replaced by an activated electrophilic ketone moiety. These analogs of the peptidase substrates provide specific enzyme inhibitors for a variety of proteases, the inhibition of which will have useful physiological consequences in a variety of disease states.

(19)



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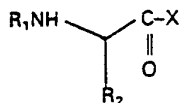
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(54) Novel peptidase inhibitors.

(57) This invention relates to analogs of peptidase substrates in which the amide group containing the scissile amide bond of the substrate peptide has been replaced by an activated electrophilic ketone moiety.

Their general formula is



the hydrates thereof, and the pharmaceutically acceptable salts thereof wherein

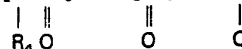
R₁ may be hydrogen, an amino protecting group, an α-amino acid or a peptide, each of said α-amino acid or peptide optionally bearing an amino protecting group

R₂ is the R group side chain of an α-amino acid building block,

X is X₁ or X₂ wherein

X₁ is -CF₃, -CF₂H, CO₂R₃ or -CONHR₃,

X₂ is -CF₂CHCR₅Y, -CF₂CR₅Y or -CR₅Y,



R₃ is hydrogen, alkyl, phenyl, benzyl, cyclohexyl or cyclohexylmethyl,

R₄ is the R group side chain of an α-amino acid building block,

R₅ is an α-amino acid or peptide building block, or deleted

Y is -NHR₃ or -OR₃, said α-amino building blocks being selected from Groups A, B, C, D, E, F, G, K and J, members of these Groups being

EP 0 195 212 A3



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EUROPEAN SEARCH REPORT

Page 195212

Application Number

EP 86 10 1437

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl. 4)
P,X	JOURNAL OF MEDICINAL CHEMISTRY, vol. 28, no. 11, November 1985, pages 1553-1555, American Chemical Society, Washington, D.C., US; S. THAISRIVONGS et al.: "Difluorostatine-and difluorostatone-containing peptides as potent and specific renin inhibitors" * Whole article, especially page 1554, table 1 *	1,30-31	
A,P	BIOCHEMISTRY, vol. 24, no. 8, 9th April 1985, pages 1813-1817, American Chemical Society, Washington, D.C., US; M.H. GELB et al.: "Fluoro ketone inhibitors of hydrolytic enzymes" * Page 1814, table 1; page 1815, tables 2,3; page 1816, table 4 *	1,36-43	
E	EP-A-0 189 305 (ICI) * Whole document, especially pages 65-66, example 2; pages 240-241, claim 1 *	1-55	
The present search report has been drawn up for all claims			TECHNICAL FIELDS SEARCHED (Int. Cl.4)
Place of search THE HAGUE		Date of completion of the search 26-06-1988	Examiner MASTURZO P.
CATEGORY OF CITED DOCUMENTS X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document I : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document			

VOSSIUS & PARTNER

PATENTANWÄLTE
EUROPEAN PATENT ATTORNEYS

0195212

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Case: C-33,761 EP
MERRELL DOW PHARMACEUTICALS INC.

February 4, 1986

NOVEL PEPTIDASE INHIBITORS

This invention relates to protease enzyme inhibitors useful for a variety of physiological end-use applications.

5 In its broad aspects, this invention relates to analogs of peptidase substrates in which the amide group containing the scissile amide bond of the substrate peptide has been replaced by an activated electrophilic ketone moiety such as fluoromethylene ketone or α -keto
10 carboxyl derivatives. These analogs of the peptidase substrates provide specific enzyme inhibitors for a variety of proteases, the inhibition of which will have useful physiological consequences in a variety of disease states.

15 In its more specific aspects, this invention relates to activated electrophilic ketone derivatives of certain peptidase substrates which are useful in inhibiting serine-, thiol-, carboxylic acid- and metallo- dependent
20 protease enzymes, the inhibition of which will have useful physiological consequences in a variety of disease states.

Still more specifically, this invention relates to activated electrophilic ketone derivatives of peptidase substrates which fall within the following generic groupings characterized according to their active site dependencies. Such generic groupings are:

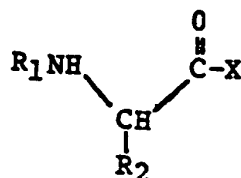
I. Serine Dependent Enzymes: These include such enzymes as Elastase (human leukocyte), Cathepsin G, Thrombin, Plasmin, C-1 Esterase, C-3-Convertase, Urokinase, Plasminogen activator, Acrosin, β -Lactamase, D-Alanine-D-Alanine Carboxypeptidase, Chymotrypsin, Trypsin and kallikreins.

II. Thiol Dependent Enzymes: Cathepsin B.

III. Carboxylic Acid Dependent Enzymes: These include such specific enzymes as Renin, Pepsin and Cathepsin D.

IV. Metallo Dependent Enzymes: These include Angiotensine Converting Enzyme, Enkephalinase, Pseudomonas Elastase and Leucine Aminopeptidase.

The contemplated peptidase inhibitors of the foregoing enzymes are selected from the generic formula



I

including the hydrates thereof, and the pharmaceutically acceptable salts thereof wherein X embraces subgroups X_1 and X_2 ,

wherein X_1 is $-CF_2H$, $-CF_3$, CO_2R_3 or $-CONHR_3$, and
 X_2 is $-CF_2\overset{\overset{R_4}{\parallel}}{CH}CR_5Y$, $-CF_2\overset{\overset{O}{\parallel}}{C}R_5Y$ and $-\overset{\overset{O}{\parallel}}{C}R_5Y$,

5 R_2 is the "R group" side chain of the α -amino acid building block responsible for directing the inhibitor to the active site of the enzyme,

R_1 may be hydrogen, an amino protecting group selected from Group K, an α -amino acid or a peptide comprised of a number of α -amino acid building blocks, each of
 10 said α -amino acid or peptide optionally bearing an amino protecting group preferably selected from Group K,

R_3 may be H, C₁-4 straight or branched alkyl, phenyl, cyclohexyl, cyclohexylmethyl or benzyl,

15 R_4 is the specific R-group side chain of an α -amino acid building block for that peptidase substrate analog,

R_5 is an α -amino acid or peptide building block, or is deleted (sometimes herein stated "or is zero") and

Y is NHR_3 or OR_3 ,

20 Unless otherwise stated the α -amino acid building blocks of these peptidase substrates are preferably in their L-configuration.

Before further defining and/or illustrating the scope of the peptidase substrate inhibitors embraced by formula
 25 I, it may be convenient to state some of the more basic concepts related to peptides. For example, except for proline, all of the α -amino acids found in proteins have, as a common denominator, a free carboxyl group and a free unsubstituted amino group on the α -carbon atom (in pro-
 30 line, since proline's α -amino group is substituted it is really an α -imino acid, but for convenience, it will also be spoken of as an α -amino group). Additionally each α -amino acid has a characteristic "R-group", the R-group being the side-chain, or residue, attached to the α -carbon

atom of the α -amino acid. For example, the R-group side chain for glycine is hydrogen, for alanine it is methyl, for valine it would be isopropyl. (Thus, throughout this specification the R₂ or R₄ moiety is the side-chain

5 R-group for each indicated α -amino acid). For the specific R-groups - or side chains - of the α -amino acids reference to A. L. Lehninger's text on Biochemistry (see particularly Chapter 4) would be helpful.

As a further convenience for defining the scope of
 10 the compounds embraced by the generic concept of Formula I, as well as the sub-generic concepts relating to each of the individual enzymes involved in this invention, various α -amino acids have been classified into a variety of groups which impart similar functional characteristics for
 15 each of the specific enzymes to be inhibited by the peptidase substrates of Formula I. These groups are set forth in Table II and the recognized abbreviations for the α -amino acid blocks are set forth in Table I.

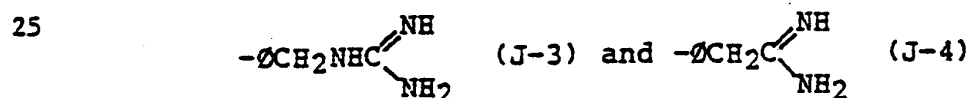
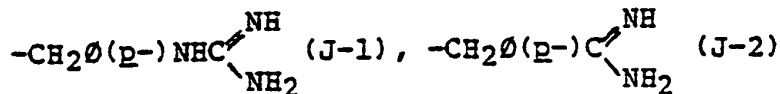
TABLE I

20	<u>Amino Acid</u>	<u>Symbol</u>
	Alanine	Ala
	Arginine	Arg
	Asparagine	Asn
	Aspartic acid	Asp
25	Asn + Asp	Asx
	Cysteine	Cys
	Glutamine	Gln
	Glutamic acid	Glu
	Gln + Glu	Glx
30	Glycine	Gly
	Histidine	His
	Isoleucine	Ile
	Leucine	Leu
	Lysine	Lys
35	Methionine	Met
	Phenylalanine	Phe
	Proline	Pro

	Serine	Ser
	Threonine	Thr
	Tryptophan	Trp
	Tyrosine	Tyr
5	Valine	Val
	Norvaline	n-Val
	n-Leucine	n-Leu
	1-Naphthylalanine	Nal(1)
	2-Indolinecarboxylic	
10	Acid	Ind
	Sarcosin	Sar

TABLE II

- Group A: Lys and Arg
 15 B: Glu, Asp
 C: Ser, Thr, Gln, Asn, Cys, His
 D: Pro, Ind
 E: Ala, Leu, Ile, Val, n-Val, Met, n-Leu and N-methyl derivatives
 F: Phe, Tyr, Trp, Nal(1), and N-methyl derivatives
 20 G: Gly, Sar
 J:

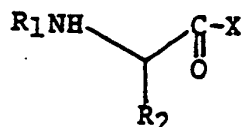


with O, of course, representing phenyl

- K: Acetyl (Ac), Succinyl (Suc), Benzoyl (Bz), t-
 30 Butyloxycarbonyl (Boc), Carbobenzoxy (CBZ), Tosyl
 (Ts), Dansyl (DNS), Isovaleryl (Iva),
 Methoxysuccinyl (MeOSuc), 1-Adamantanesulphonyl
 (AdSO₂), 1-Adamantaneacetyl (AcAc), 2-
 35 Carboxybenzoyl (2-CBZ) and such other terminal
 amino protecting groups which are functionally
 equivalent thereto.

In light of the foregoing, the defined compounds of
 formula I may also be stated as being:

An activated electrophilic ketone-bearing peptidase
 40 inhibitor of the formula



the hydrates thereof, and the pharmaceutically acceptable
 5 salts thereof wherein

R₁ may be hydrogen, an amino protecting group selected
 from Group K, and α-amino acid or a peptide comprised
 of a number of α-amino acid building blocks, each of
 said α-amino acid or peptide optionally bearing an
 10 amino protecting group selected from Group K,

R₂ is the R group side chain of an α-amino acid building
 block,

X is X₁ or X₂ wherein

X₁ is CF₃, CF₂H, CO₂R₃ or -CONHR₃,

15 X₂ is $-\text{CF}_2\underset{\text{R}_4\text{O}}{\underset{\parallel}{\text{CH}}}\text{CR}_5\text{Y}$, $-\text{CF}_2\underset{\text{O}}{\underset{\parallel}{\text{C}}}\text{R}_5\text{Y}$ or $-\underset{\text{O}}{\underset{\parallel}{\text{C}}}\text{R}_5\text{Y}$,

R₃ is hydrogen, C₁-4 straight or branched alkyl, phenyl,
 benzyl, cyclohexyl or cyclohexylmethyl,

R₄ is the R group side chain of an α-amino acid building
 20 block,

R₅ is an α-amino acid or a peptide comprised of α-amino
 acid building blocks, or is deleted,

Y is -NHR₃ or -OR₃

wherein the said α-amino acid and peptide moieties are
 25 building blocks selected from Groups A, B, C, D, E,
 F, G and J, and K is a terminal amino protecting group,
 members of these groups being

Group A: Lys and Arg,

B: Glu and Asp,

30 C: Ser, Thr, Gln, Asn, Cys and His,

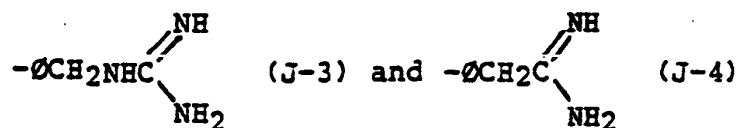
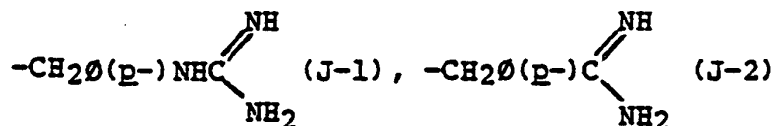
D: Pro, Ind

E: Ala, Leu, Ile, Val, n-Val, Met and n-Leu, and N-methyl derivatives,

F: Phe, Tyr, Trp, Nal(1) and N-methyl derivatives

G: Gly, Sar

J:



with Ø, of course, representing phenyl,

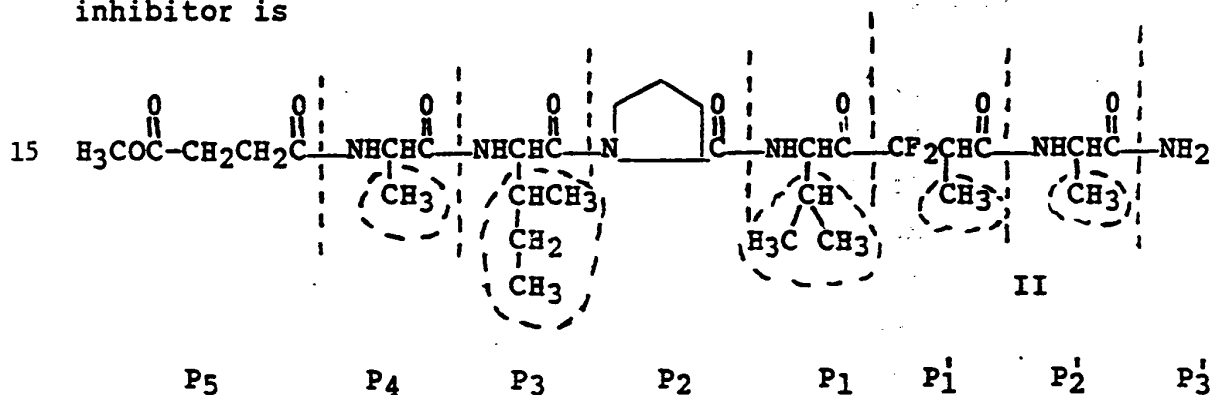
K: Acetyl (Ac), Succinyl (Suc), Benzoyl (Bz), t-Butyloxycarbonyl (Boc), Carbobenzoxy (CBZ), Tosyl (Ts), Dansyl (DNS), Isovaleryl (Iva), Methoxysuccinyl (MeOSuc), 1-Adamantanesulphonyl (AdSO₂), 1-Adamantaneacetyl (AcAc), 2-Carboxybenzoyl (2-CBZ) and such other terminal amino protecting groups which are functionally equivalent thereto.

To illustrate those compounds which are useful as enzyme inhibitors for human leukocyte elastase, and to serve as a medium of instruction for a better understanding of the scope of compounds embraced within the generic formula I, (and its sub-generic formulae for each of the involved enzymes herein disclosed) the following formula (Ia) represents the sub-generic class defining those compounds within the scope of inhibitors of human leukocyte elastase:



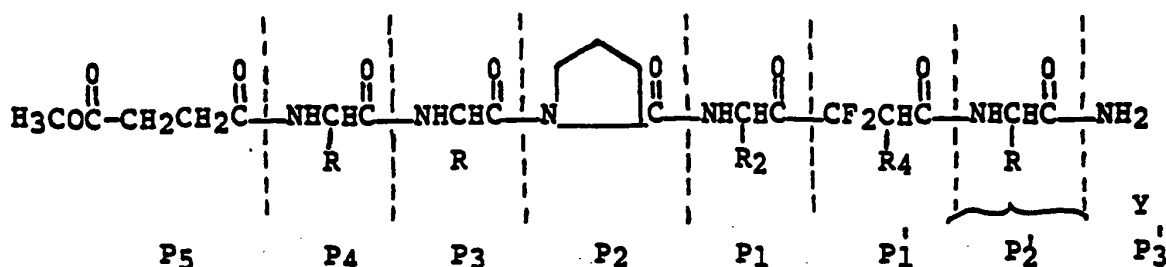
wherein R₂ is the side chain of the depicted enzyme-
 directing α-amino acid building block (P₁),
 R₁ is as previously defined (Comprised of P₂-P_n blocks),
 and
 X is the moiety conferring the electrophilic character to
 its adjacent carbonyl consisting of either X₁ or X₂
 as previously defined generically in formula I.

Still for instructional purposes, the structural
 formula for the most preferred human leukocyte elastase
 inhibitor is



20 the vertical dotted lines setting off each moiety
 constituting the specific arrangement for that particular
 peptidase inhibitor. Except for proline and 2-indoline
 carboxylic acid the moieties encircled within the dotted
 lines represent the R-group side chains of the α-amino
 25 acid building block (see pages 69-71 of the above cited
 Lehninger's text) or 1-naphthylmethyl.

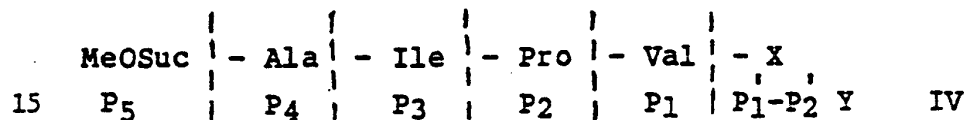
Still another way of representing the foregoing sub-
 strate is by the formula



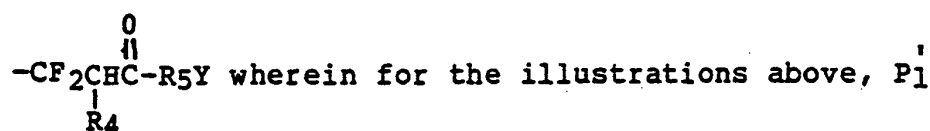
III

- 5 with R, R₂, R₄ being the side chain residues of the particular α-amino acid, the amino acid building block of P₂' (P₂ primed) being the R₅, if any, of X₂ and the terminal P₃' being a specific of the Y radical of X₂, P₅ being the terminal moiety sometimes referred generically as (P_n) and
- 10 P₂-P₃-P₄ being the remaining α-amino acid building blocks of that R₁ moiety.

Still another, and the most convenient method for simply conveying the structure involved is the formula



wherein X consists of P₁', P₂'Y when representative of



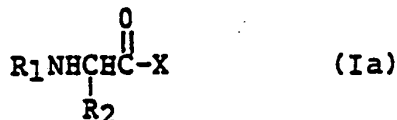
- 20 bears the -CH₃ side chain R-group for R₄ and P₂' bears the R₅ amino acid block bearing the -CH₃ side chain and Y is NH₂, and P₁-P₅ are shorthand designations for the depicted P₁-P₅ moieties of the above structures II and III.

- To expand on structure IV and (Ia) as it encompasses
- 25 the scope of the other X moieties attached to the same P₁-P₅ moieties, the following seven structures are shown:

- (a) MeOSuc-Ala-Ile-Pro-Val-CF₂H (i.e., X₁ is -CF₂H).
- (b) MeOSuc-Ala-Ile-Pro-Val-CF₃ (i.e., X₁ is -CF₃).
- (c) MeOSuc-Ala-Ile-Pro-Val-COOH (i.e., X₁ is -CO₂R₃ with R₃ being H).
- 5 (d) MeOSuc-Ala-Ile-Pro-Val-CONH₂ (i.e., X₁ is -CONHR₃ with R₃ being H).
- (e) MeOSuc-Ala-Ile-Pro-Val-CF₂CHC-NHCHC-NH₂ (i.e., X₂ is
- $$\begin{array}{c} \text{R}_4\text{O} \\ | \\ \text{CF}_2\text{CHC}-\text{R}_5\text{Y} \end{array} \quad \begin{array}{c} \text{H}_3\text{C} \\ | \\ \text{CHC} \\ || \\ \text{O} \end{array} \quad \begin{array}{c} \text{H}_3\text{C} \\ | \\ \text{CHC} \\ || \\ \text{O} \end{array}$$
- 10 CF₂CHC-R₅Y with R₄ being the side chain of alanine and R₅ is -NHCH-C- with R₅ being the side chain of
- $$\begin{array}{c} \text{R}_5 \\ | \\ \text{CH} \\ || \\ \text{O} \end{array}$$
- alanine and Y is NH₂.
- (f) MeOSuc-Ala-Ile-Pro-Val-CF₂C-Ala-NH₂ (i.e., X₂ is
- $$\begin{array}{c} \text{O} \\ || \\ \text{CF}_2\text{C}-\text{R}_5\text{Y} \end{array} \quad \begin{array}{c} \text{O} \\ || \\ \text{C} \\ || \\ \text{O} \end{array}$$
- 15 CF₂C-R₅Y with R₅ being as defined in (e) above, and Y is NH₂).
- (g) MeOSuc-Ala-Ile-Pro-Val-C-Ala NH₂ (i.e., X₂ is -C-R₅Y
- $$\begin{array}{c} \text{O} \\ || \\ \text{C} \\ || \\ \text{O} \end{array} \quad \begin{array}{c} \text{O} \\ || \\ \text{C} \\ || \\ \text{O} \end{array}$$
- wherein R₅ is as defined in (e) above and Y is NH₂.

- 20 It is also convenient when defining the substrate compounds according to the foregoing formula IV convention designation to define the -CF₂CHC moiety of X₂ as [CF₂-α-
- $$\begin{array}{c} \text{R}_4\text{O} \\ | \\ \text{CHC} \\ || \\ \text{O} \end{array}$$
- amino acid] wherein the name of the α-amino acid to which
- 25 R₄ is a side chain is indicated such as, for example, in this convention -CF₂CHC becomes [CF₂ Alal. This will
- $$\begin{array}{c} \text{H}_3\text{C} \\ | \\ \text{CHC} \\ || \\ \text{O} \end{array}$$
- facilitate the writing and comprehension of the so-defined structures.

- 30 Utilizing the foregoing illustrations those compounds of formula I which are useful as inhibitors for human leukocyte elastase are represented by the formula



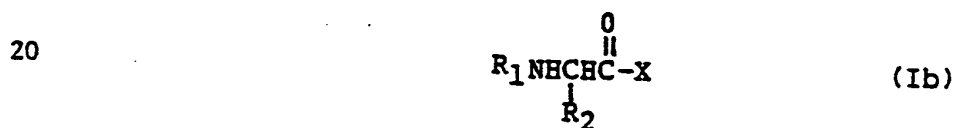
- wherein R₂ is the side chain of the α-amino acids of Groups E and G, with nor-valine and valine being preferred,
- R₁ is -P₂P₃P₄P₅ with P₂ being α-amino acid blocks of Groups D, E and F, with proline being preferred,
- P₃ is the α-amino acid blocks of Groups D or E, with isoleucine being preferred,
- P₄ is the α-amino acid blocks of Groups E or zero with alanine being preferred (when P_n is zero then that particular moiety does not appear in the structure, i.e. it is deleted),
- P₅ is the terminal moiety of Group K with methoxysuccinyl being preferred,
- X is any of the X₁ or X₂ moieties defined for formula I with R₅ being an α-amino acid block of Groups E and G with alanine being preferred and Y is NH₂, and
- R₄ is an R group side chain of Groups E and G with alanine being-preferred.

Human leukocyte elastase is released by polymorphonuclear leukocytes at sites of inflammation and thus is a contributing cause for a number of disease states. Thus the peptidase substrates of Formula (Ia) have an anti-inflammatory effect useful in the treatment of gout, rheumatoid arthritis and other inflammatory diseases, and in the treatment of emphysema. In their end-use application the enzyme inhibitory properties of the compounds of (Ia) are readily ascertained by standard biochemical techniques well known in the art. Potential dose range for their end-use application will of course depend upon the nature and severity of the disease state as determined by the attending diagnostician with the range of 0.01 to 10 mg/kg

body weight per day being useful for the aforementioned disease states. The preferred compounds for this enzyme are:

- MeOSuc-Ala-Ile-Pro-Val-[CF₂-Ala]Ala-NH₂,
 MeOSuc-Ala-Ile-Pro-Val-CF₃,
 5 MeOSuc-Ala-Ile-Pro-Val-CO₂Me,
 MeOSuc-Ala-Ile-Pro-Val-CF₂COOEt,
 MeOSuc-Ala-Ile-Pro-Val-CHF₂,
 MeOSuc-Ala-Ala-Pro-Val-CO₂Me,
 MeOSuc-Ala-Ala-Pro-Val-[CF₂-Ala]Ala-NH₂,
 10 MeOSuc-Ala-Ala-Pro-Val-CF₃,
 [αN-(AdSO₂)]-[EN-(2-CBz)]-Lys-Pro-Val-[CF₂Ala]Ala-NH₂
 [αN-(AdSO₂)]-[EN-(2-CBz)]-Lys-Pro-Val-CHF₂
 [αN-(AdSO₂)]-[EN-(2-CBz)]-Lys-Pro-Val-CO₂Me
 MeOSuc-Ala-Ile-Pro-Val-CO₂Me,
 15 MeOSuc-Ala-Ile-Pro-Val-CO₂H.

Those compounds of Formula I which are useful as inhibitors of Cathepsin G are represented by the structural formula



wherein X₁, X₂, R₃, R₄, R₅ and Y are as defined for human leukocyte elastase (Formula Ia),

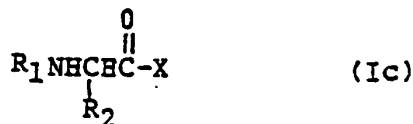
- R₁ is P₂-P₃-P₄-P₅ with P₂ being selected from Groups D, E,
 25 G or K with proline or benzoyl being preferred,
 P₃ is selected from Groups E or G with alanine being preferred,
 P₄ is selected from Groups E, G or is deleted with alanine being preferred,
 30 P₅ is selected from Group K with succinyl being preferred.

R₂ is selected from Groups E and F but preferably is Phe side chain.

The end-use application of the compounds (Ib) inhibiting Cathepsin G is the same as for human leukocyte inhibitors, including arthritis, gout and emphysema, but also embracing the treatment of glomerulonephritis and lung infestations caused by infections in the lung. For their end-use application, the potency and other biochemical parameters of the enzyme inhibiting characteristics of the compounds of (Ib) are readily ascertained by standard biochemical techniques well known in the art. Actual dose ranges for their specific end-use application will, of course, depend upon the nature and severity of the disease state of the patient or animal to be treated as determined by the attending diagnostician. It is to be expected that the general end-use application dose range will be about 0.01 to 10 mg per kg per day for an effective therapeutic effect. Preferred compounds of formula Ib are:

Suc-Ala-Ala-Pro-Phe-X₁, and specifically
 Suc-Ala-Ala-Pro-Phe-CF₃,
 Suc-Ala-Ala-Pro-Phe-COOH, and
 Suc-Ala-Ala-Pro-Phe-COOMe,
 Suc-Ala-Ala-Pro-Phe-CF₂H,
 Suc-Ala-Ala-Pro-Phe[CF₂Ala]OH,
 Suc-Ala-Ala-Pro-Phe-CF₂-COOEt

Those compounds of Formula I which are useful as inhibitors of thrombin are represented by the formula



0195212
wherein X is X₁ or X₂ as defined in formula I with Y being

OH,

- R₅ is preferably the glycine amino acid block or is a member of Group E or D or is zero,
- 5 R₄ is selected from Group C or G but preferably is a glycine or serine side chain,
- R₂ is preferably the arginine side chain but may also be selected from Groups A and J,
- R₁ is (a)-P₂-P₃, (b)-P₂ or (c)-P₂-P₃-P₄ with
- 10 (a) P₂ is selected from Groups E or F, preferably proline, P₃ is selected from Group F, each P₃ being in the D - configuration preferably D-Phe,
- (b) P₂ is selected from Group K but preferably is dansyl or tosyl,
- 15 (c) P₂ is selected from Group E but preferably is alanine, P₃ is selected from Groups G and E but preferably is serine, P₄ is selected from Groups G and E or is zero but preferably is Phe.

The compounds embraced by Formula (Ic) inhibit thrombin and therefore, as in the use of heparin, the compounds may be used as the initial anticoagulant agent in thrombophlebitis and coronary thrombosis. For their end-use application, the potency and other biochemical parameters of the enzyme inhibiting characteristics of the compounds of (Ic) are readily ascertained by standard biochemical techniques well known in the art. Actual dose ranges for their specific end-use application will, of course, depend upon the nature and severity of the disease state of the patient or animal to be treated as determined by the attending diagnostician. It is to be expected that the general end-use application dose range will be about 0.01 to 10 mg per kg per day for an effective therapeutic effect. Preferred compounds are as expressed for Cathepsin G and also include:

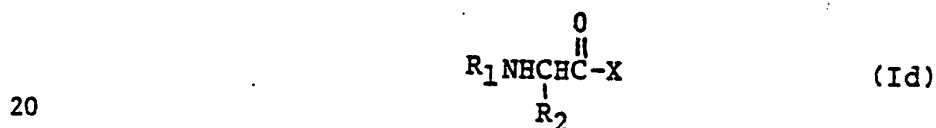
20

25

30

- H-(D)-Phe-Pro-Arg-CF₃,
 H-(D)-Phe-Pro-Arg-COOH,
 H-(D)-Phe-Pro-Arg-COO-n-butyl,
 DNS-Arg-CF₃,
 5 DNS-Arg-COOH,,
 DNS-Arg-COO-n-butyl,
 H-Phe-Ser-Ala-CF₃,
 H-Phe-Ser-Ala-COOH,
 H-Phe-Ser-Ala-COO-n-butyl,
 10 $\text{P}-[\text{H}_2\text{NC}(\text{NH})\text{NH}]-\text{C}_6\text{H}_4-\text{CH}_2\text{CH}(\text{NHBz})\overset{\text{O}}{\underset{\text{O}}{\text{C}}}\text{CF}_3$, $\text{P}-[\text{H}_2\text{NC}(\text{NH})\text{NH}]-\text{C}_6\text{H}_4-\text{CH}_2\text{CH}(\text{NHBz})\overset{\text{O}}{\underset{\text{O}}{\text{C}}}\text{CF}_3$
 $\text{P}-[\text{H}_2\text{NC}(\text{NH})\text{NH}]-\text{C}_6\text{H}_4-\text{CH}_2\text{CH}(\text{NHBz})\overset{\text{O}}{\underset{\text{O}}{\text{C}}}[\text{CF}_2\text{Gly}]\text{Pro-OH}$,
 $\text{P}-[\text{H}_2\text{NC}(\text{NH})\text{NH}]\text{C}_6\text{H}_4\text{CH}_2\text{CH}(\text{NHBz})\text{COOH}$

- 15 The compounds of Formula I which are useful as inhibitors of chymotrypsin are represented by the structural formula



- wherein X₁, X₂, R₃, R₄ R₅ and Y are as defined for compounds of Ia, and R₁ is -P₂-P₃-P₄-P₅ with,
 P₂ being selected from Groups D, E, G or K with benzoyl being preferred,
 25 P₃ is selected from Groups E or G or is zero with alanine being preferred,
 P₄ is selected from Groups E or G or is deleted with alanine being preferred,
 P₅ is selected from Group K with succinyl being preferred,
 30 or is zero when P₂ is K, and
 R₂ is selected from Groups E and F but preferably is Phe or Tyr side chains.

The end-use application of the compounds (Id) inhibiting chymotrypsin is in the treatment of pancreatitis. For their end-use application, the potency and other biochemical parameters of the enzyme inhibiting characteristics of the compounds of (Id) are readily ascertained by standard biochemical techniques well known in the art. Actual dose ranges for their specific end-use application will, of course, depend upon the nature and severity of the disease state of the patient or animal to be treated as determined by the attending diagnostician. It is to be expected that the general end-use application dose range will be about 0.01 to 10 mg per kg per day for an effective therapeutic effect. Preferred compounds are as expressed for Cathepsin G and also include:

15 Bz-Phe-CF₃,
 Bz-Phe-COOH,
 Bz-Phe-COOME,
 Bz-Tyr-CF₃,
 Bz-Tyr-COOH,
 20 Bz-Tyr-COOME,
 Bz-Phe-CHF₂,
 Bz-Phe-CF₂COOEt
 Bz-Phe-[CF₂-Gly]Gly-OH

The compounds of Formula I which are useful as inhibitors of trypsin are represented by the structural formula



30 wherein X is X₁ or X₂ as defined in formula I with Y being OH,

- R₅ is selected from Groups G, E or D or is zero but preferably is glycine,
- R₄ is an R group side chain of Groups C or G but preferably is glycine or serine side chain,
- 5 R₂ is selected from Groups A or J but preferably is the arginine side chain,
- R₁ is selected from (a)-P₂-P₃, (b)-P₂ or (c)-P₂-P₃-P₄ with
- (a) P₂ is selected from Groups E or F but is preferably proline or alanine, P₃ is selected
- 10 from Group F, (each being in the D configuration) but preferably is D -Phe,
- (b) P₂ is selected from Group K but preferably is dansyl or tosyl,
- (c) P₂ is selected from Group D or E but preferably
- 15 is proline or alanine, P₃ is selected from Groups G and E but preferably is serine, P₄ is selected from Groups G and E or is zero but preferably is Phe.

The end-use application of the compounds (Ie) inhibi-

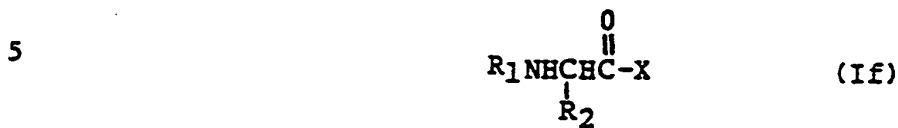
20 ting trypsin is in the treatment of pancreatitis. For their end-use application, the potency and other biochemical parameters of the enzyme inhibiting characteristics of the compounds of (Ie) are readily ascertained by standard biochemical techniques well known in the art. Actual

25 dose ranges for their specific end-use application will, of course, depend upon the nature and severity of the disease state of the patient or animal to be treated as determined by the attending diagnostician. It is to be expected that the general end-use application dose range

30 will be about 0.01 to 10 mg per kg per day for an effective therapeutic effect. The preferred compounds useful for inhibiting trypsin are the same as for the inhibitors of thrombin.

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The compounds of Formula I which are useful as inhibitors of plasmin are represented by the structural formula



wherein X is X₁ or X₂, with CF₃, COOH, COOMe, and CF₂COOEt being preferred,

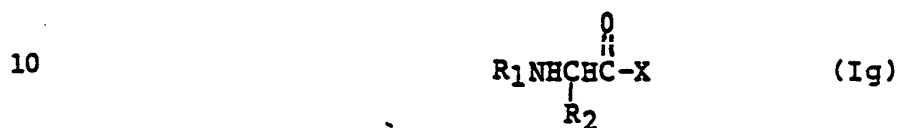
10 R₁ is -P₂-P₃-P₄ with P₂ being selected from Group F but preferably is Phe, P₃ is selected from Groups B or F but preferably is Glu, and P₄ is selected from Group K but preferably is dansyl,

R₂ is selected from Groups A and J but preferably is the lysine side chain.

15 The compounds embraced by formula (If) inhibit plasmin and are therefore antiproliferative agents useful in treating excessive cell growth, particularly in the treatment of benign prostatic hypertrophy and prostatic carcinoma, and in the treatment of psoriasis. For their
20 end-use application, the potency and other biochemical parameters of the enzyme inhibiting characteristics of the compounds of (If) are readily ascertained by standard biochemical techniques well known in the art. Actual dose ranges for their specific end-use application will, of
25 course, depend upon the nature and severity of the disease state of the patient or animal to be treated as determined by the attending diagnostician. It is to be expected that the general end-use application dose range will be about 0.01 to 10 mg per kg per day for an effective therapeutic
30 effect. The preferred compounds are:

DNS-Glu-Phe-Lys-CHF₂
 DNS-Glu-Phe-Lys-COOH
 DNS-Glu-Phe-Lys-CF₃
 DNS-Glu-Phe-Lys-COOMe
 5 DNS-Gly-Phe-Lys-CF₂COOEt

The compounds of Formula I which are useful as inhibitors of C₁-esterase are represented by the structural formula



wherein X generically is X₁ or X₂ with X₁ being preferred particularly when X₁ is CO₂R₃ or -CF₃,
 R₂ is selected from Groups A and J but preferably is Arg,
 15 R₁ generically is -P₂-P₃ with P₂ being selected from Groups E, G, D, C, F, A or B with Ala. being preferred, and P₃ is selected from Group K with CBZ being preferred,
 R₄ is selected from Group E,
 20 R₅ is selected from Group E and Y is preferably NH₂.

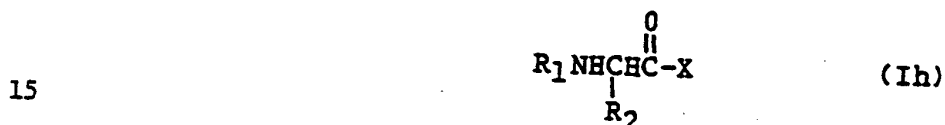
The compounds embraced by formula (Ig) inhibit C₁-esterase and are therefore useful in treating systemic lupus, arthritis, autoimmune hemolytic anemia and glomerulonephritis. For their end-use application, the
 25 potency and other biochemical parameters of the enzyme inhibiting characteristics of the compounds of (Ig) are readily ascertained by standard biochemical techniques well known in the art. Actual dose ranges for their specific end-use application will, of course, depend upon
 30 the nature and severity of the disease state of the patient or animal to be treated as determined by the

attending diagnostician. It is to be expected that the general end-use application dose range will be about 0.01 to 10 mg per kg per day for an effective therapeutic effect. The preferred compounds are:

- 5 CBZ-Ala-Arg-CF₃,
 CBZ-Ala-Arg-COOH,
 CBZ-Ala-Arg-COOMe,
 CBZ-Ala-(p-gua)*-Phe-CF₂COOEt,
 CBZ-Ala-(p-gua)-Phe[CF₂Ala]NH₂,

10 *gua is guanidino

The compounds of Formula I which are useful as inhibitors of C₃-convertase are represented by the formula



wherein X generically is X₁ or X₂ with X₂ being preferred, R₄ preferably being the alanine side chain, but is also Group E,

- R₅ is zero and Y is OR₃ (i.e., R₅Y is OR₃),
 20 R₂ is selected from Groups A or J, with Arg being preferred,
 R₁ is -P₂-P₃-P₄ with P₂ being selected from Groups E or F, with Ala being preferred, P₃ is selected from Groups E or F with Leu being preferred, and
 25 P₄ is selected from Group K with Bz being preferred.

The compounds embraced by formula (Ih) inhibit C₃-convertase and are therefore useful in treating systemic lupus, arthritis, autoimmune hemolytic anemia and glomerulonephritis. For their end-use application, the
 30 potency and other biochemical parameters of the enzyme inhibiting characteristics of the compounds of (Ih) are

readily ascertained by standard biochemical techniques well known in the art. Actual dose ranges for their specific end-use application will, of course, depend upon the nature and severity of the disease state of the patient or animal to be treated as determined by the attending diagnostician. It is to be expected that the general end-use application dose range will be about 0.01 to 10 mg per kg per day for an effective therapeutic effect. The preferred compounds are:

- 10 Bz-Leu-Ala-Arg-CF₃,
 Bz-Leu-Ala-Arg-CHF₂,
 Bz-Leu-Ala-Arg-CF₂-COO-CH₂Ø
 Bz-Leu-Ala-Arg[CF₂-Ala]OCH₂Ø
 Bz-Leu-Ala-Arg-COOCH₂Ø.

- 15 The compounds of formula I which are useful as inhibitors of Urokinase are represented by the formula



- 20 wherein X generically is X₁ or X₂ with X₁ being preferred and CO₂R₃ and -CF₃ being most preferred,
 R₄ is Group E,
 R₅ is Group E, and
 Y is NH₂,
 25 R₁ generically is -P₂-P₃ with P₂ being selected from Groups E and G with Ala and Gly being preferred, and P₃ is selected from Group B with Glu being preferred,
 R₂ is selected from Groups A and J with Arg being preferred.

- 30 Preferred Urokinase inhibitors are:

K-Glu-Gly-Arg-CF₂H,
 K-Glu-Gly-Arg-CF₃,
 K-Glu-Gly-Arg-COOH,
 K-Glu-Gly-Arg-CONH₂,
 5 K-Glu-Gly-(p-gua)*Phe-[CF₂Ala]-Ala-NH₂, and
 K-Gly-Gly(p-gua)*Phe-CF₂CONH₂,

** (p-gua) being para-guanidino

The compounds of formula (Ii) inhibit urokinase and therefore are useful in treating excessive cell growth
 10 disease states. As such the compounds are useful in the treatment of benign prostatic hypertrophy and prostatic carcinoma, the treatment of psoriasis, and in their use as abortifacients. For their end-use application, the potency and other biochemical parameters of the enzyme
 15 inhibiting characteristics of the compounds of (Ii) are readily ascertained by standard biochemical techniques well known in the art. Actual dose ranges for their specific end-use application will, of course, depend upon the nature and severity of the disease state of the
 20 patient or animal to be treated as determined by the attending diagnostician. It is to be expected that the general end-use application dose range will be about 0.01 to 10 mg per kg per day for an effective therapeutic effect.

25 The compounds of Formula I which are useful as inhibitors of plasminogen activator are represented by the structural formula



wherein X generically is X₁ or X₂ with X₁ being preferred
and -CF₃, COOH and COOMe being most preferred,

R₄ is Group E,

R₅ is Group E,

5 Y is NH₂ when X is X₂,

R₁ generically is -P₂-P₃-P₄ wherein P₂ is Gly, P₃ is
selected from Group B with Glu being preferred, and
P₄ preferably is dansyl but also selected from Group
K, and

10 R₂ is selected from Groups A and J with Arg being
preferred.

Preferred compounds are:

DNS-Glu-Gly-Arg-COOMe,

DNS-Glu-Gly-Arg-CF₃,

15 DNS-Glu-Gly-Arg-COOH,

DNS-Glu-Gly-(p-gua)Phe-CHF₂,

DNS-Glu-Glu-(p-gua)Phe[CF₂Ala]AlaNH₂,

DNS-Glu-Gly-(p-gua)PheCF₂COOEt.

20 The compounds of the Formula (Ij) inhibit plasminogen
activator and therefore are useful in treating excessive
cell growth disease states such, for example, being useful
in the treatment of benign prostatic hypertrophy and pro-
static carcinoma, in the treatment of psoriasis and in
their use as abortifacients. For their end-use applica-
25 tion, the potency and other biochemical parameters of the
enzyme inhibiting characteristics of the compounds of (Ij)
are readily ascertained by standard biochemical techniques
well known in the art. Actual dose ranges for their spe-
cific end-use application will, of course, depend upon the
30 nature and severity of the disease state of the patient or
animal to be treated as determined by the attending diag-
nostician. It is to be expected that the general end-use
application dose range will be about 0.01 to 10 mg per kg
per day for an effective therapeutic effect.

The compounds of Formula I which are useful as inhibitors of acrosin are represented by the structural formula



wherein X generically is X₁ or X₂, with X₁ being preferred especially when X₁ is -CF₃, CHF₂, COOH or COOMe.

10 When X is X₂, R₄ is Group E, R₅ is Group E, or is deleted and Y is NH₂,

R₁ generically is -P₂-P₃-P₄ with P₂ being selected from Group E with Leu being preferred, P₃ is selected from Group E with Leu being preferred, P₄ is selected from Group K with Boc being preferred.

15 R₂ is selected from Groups A and J with Arg being preferred. Preferred compounds are:

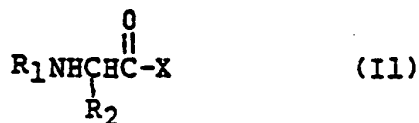
Boc-Leu-Leu-Arg-CF₂H,
 Boc-Leu-Leu-Arg-CF₃,
 Boc-Leu-Leu-Arg-COOH,
 20 Boc-Leu-Leu-(p-gua)Phe-[CF₂Ala]AlaNH₂, and
 Boc-Leu-Leu-(p-gua)PheCF₂CONH₂.

The compounds of formula (Ik) are acrosin inhibitors and therefore are useful as anti-fertility agents in that they possess the characteristics of preventing sperm from
 25 penetrating an otherwise fertilizable egg. For their end-use application, the potency and other biochemical parameters of the enzyme inhibiting characteristics of the compounds of (Ik) are readily ascertained by standard biochemical techniques well known in the art. Actual dose

ranges for their specific end-use application will, of course, depend upon the state of the patient or animal to be treated as determined by the attending diagnostician. It is to be expected that the general end-use application
 5 dose range will be about 0.01 to 10 mg per kg per day for an effective therapeutic effect.

The compounds of Formula I which are useful as inhibitors of β -lactamase are represented by the structural formula

10



with the proviso that the depicted carbonyl moiety (attached to X) may exist in its

15

chemically reduced form, (i.e., $\text{R}_1\text{NHCH}(\text{OH})\text{CH}(\text{R}_2)\text{X}$) with the

$$\begin{array}{c} \text{OH} \\ | \\ \text{R}_1\text{NHCH}-\text{CH}-\text{X} \\ | \\ \text{R}_2 \end{array}$$

reduced form being preferred,

wherein X is X_1 or X_2 with $-\text{CF}_3$, COOH and COOMe being most preferred, and R_5 is deleted when X is X_2

20

R_1 generically is P_2 , P_2 being selected from Group K with $\text{COCH}_2\emptyset$ and Bz being preferred, when X is generically X_2 ,

R_2 is selected from Groups E, G and C with glycine being preferred. The preferred compounds are:

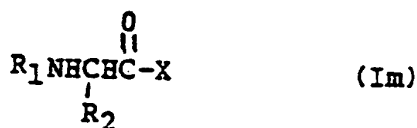
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- $\emptyset\text{CH}_2\text{COHNCH}_2\text{COCF}_3$,
 $\emptyset\text{CH}_2\text{COHNCH}_2\text{COCOCH}_3$,
 $\emptyset\text{CH}_2\text{COHNCH}_2\text{COCOCH}_2\text{CH}_3$,
 $\emptyset\text{CH}_2\text{COHNCH}_2\text{CHOHCF}_3$,
 $\emptyset\text{CH}_2\text{COHNCH}_2\text{CHOHCOCH}_3$,
 $\emptyset\text{CH}_2\text{COHNCH}_2\text{CHOHCOCH}_2\text{CH}_3$,
 $\emptyset\text{CH}_2\text{COHNCH}_2\text{COCHF}_2$,
 $\emptyset\text{CH}_2\text{COHNCH}_2\text{CHOHCF}_2\text{COOEt}$.

30

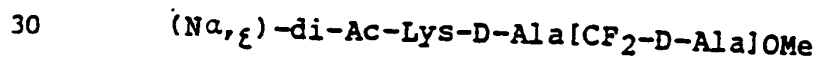
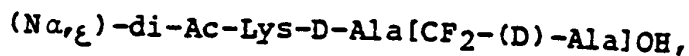
The compounds embraced by formula (II) inhibit β -Lactamase and therefore are useful in the potentiation of antibacterial agents, particularly the β -lactam antibacterials. For their end-use application, the potency and other biochemical parameters of the enzyme inhibiting characteristics of the compounds of (II) are readily ascertained by standard biochemical techniques well known in the art. Actual dose ranges for their specific end-use application will, of course, depend upon the nature and severity of the disease state of the patient or animal to be treated as determined by the attending diagnostician. It is to be expected that the general end-use application dose range will be about 0.01 to 10 mg per kg per day for an effective therapeutic effect.

The compounds of Formula I which are useful as inhibitors of D-Ala-D-Ala Carboxypeptidase



wherein X generically is X_1 or X_2 wherein when X is X_2 , R_4 is D-Ala and R_5 is deleted and Y is OH or OR_3 , R_2 is D-Ala,

R_1 generically is P_2 - P_3 with P_2 being $\text{Ac}(\text{N}\alpha, \epsilon\text{-Ac})\text{Lys}$ or Groups E and C with $\text{Ac}(\text{N}\alpha, \epsilon\text{-Ac})\text{Lys}$ being preferred, P_3 generically is selected from Group K with Ac being preferred. The preferred compounds are:

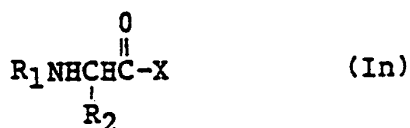


(N α , ϵ)-di-Ac-Lys-D-Ala-CHF₂,
 (N α , ϵ)-di-Ac-Lys, D-Ala-CF₂COOEt and
 N α , ϵ -di-Ac-Lys-D-AlaCF₃

The compounds embraced by formula (Im) are antibac-
 5 terial agents particularly useful against gram negative
 organisms. For their end-use application, the potency and
 other biochemical parameters of the enzyme inhibiting
 characteristics of the compounds of (Im) are readily ascer-
 10 tained by standard biochemical techniques well known in
 the art. Actual dose ranges for their specific end-use
 application will, of course, depend upon the nature and
 severity of the disease state of the patient or animal to
 be treated as determined by the attending diagnostician.
 It is to be expected that the general end-use application
 15 dose range will be about 0.01 to 10 mg per kg per day for
 an effective therapeutic effect.

The compounds of Formula I which are useful as inhi-
 bitors of Cathepsin B are represented by the structural
 formula

20



wherein X generically is X₁ or X₂ with X₁ being preferred
 and CF₃ and COOH being particularly preferred and
 25 when X is X₂, R₄ is selected from Group E with Leu
 being preferred,
 R₅ is selected from Groups G, E or F with Gly being
 preferred and Y is OH,
 R₁ generically is (a)-P₂-P₃ or (b)-P₂-P₃-P₄ wherein for

(a) P₂ is selected from Groups E and F with Phe being preferred and P₃ is selected from Group K with CBZ being preferred, and

(b) P₂ is selected from Groups E and F with Leu being preferred, -P₃ being selected from groups E and F with Leu being preferred and P₄ is selected from Group K with Ac being preferred,

R₂ is selected from Group A and J or ThrCOCH₂O, with Arg being preferred. The preferred compounds are:

10 Ac-Leu-Leu-Arg[CF₂-Leu]Gly-OH,
CBZ-Phe-Arg[CF₂-Leu]Gly-OH, and
CBZ-Phe-Thr[CF₂-Leu]Gly-OH.

OBz

CBZ-Phe-Thr-CHF₂

OBz

15 CBZ-Phe-Thr-CF₃

OBz

CBX-Phe-Thr-CF₂-CO-Gly-OH

OBz

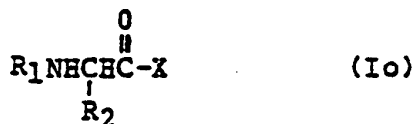
20 The compounds of formula (In) inhibit Cathepsin B and therefore are useful in treating excessive cell growth disease states such as, for example, being useful in treating benign prostate hypertrophy, prostatic carcinoma, in treating psoriasis and in their use as abortifacients.

25 For their end-use application, the potency and other biochemical parameters of the enzyme inhibiting characteristics of the compounds of (In) are readily ascertained by standard biochemical techniques well known in the art. Actual dose ranges for their specific end-use application

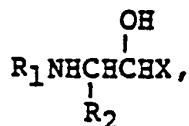
30 will, of course, depend upon the nature and severity of the disease state of the patient or animal to be treated as determined by the attending diagnostician. It is to be expected that the general end-use application dose range

will be about 0.01 to 10 mg per kg per day for an effective therapeutic effect.

The compounds of Formula I which are useful as inhibitors of renin are represented by the structural
5 formula



with the proviso that the depicted carbonyl moiety
10 attached to X may exist in its chemically reduced form, i.e.



wherein X generically is X_1 or X_2 , which when X is X_1 CF_3 ,
15 COOH or COOMe are preferred and when X is X_2 ,

R_4 is selected from Groups E, F or G with Val being preferred,

R_5 generically is $-P_2^1-P_3^1-P_4^1$, P_2^1 being from Groups E, F or
20 is deleted, with P_3^1 being selected from Groups E or F or is deleted with Ile being preferred and P_4^1 being selected from Groups E, C or F or is deleted with His being preferred and Y is OH or NH_2 ,

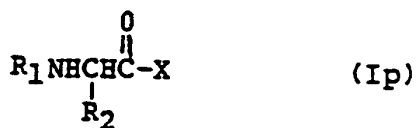
R_2 is selected from Groups E or F or is cyclohexylmethylene with Leu being preferred,

25 R_1 generically is $-P_2-P_3-P_4-P_5-P_6$ wherein P_2 is selected from Groups E, C or F with His being preferred, P_3 is selected from Groups E or F with Phe being preferred, P_4 is selected from Groups E, D, F or is deleted with Pro being preferred, P_5 is selected from
30 Groups E, C, F or is deleted with His being preferred, and P_6 is selected from Group K with MeOSuc being preferred. The preferred compounds are:

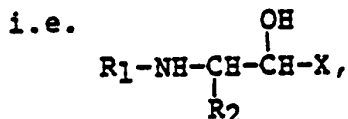
CBZ-Nal(1)-His-Leu-CHF₂,
 CBZ-Nal(1)-His-Leu-CF₃,
 CBZ-Nal(1)-His-Leu-CF₂-COOEt,
 MeOSuc-His-Pro-Phe-His-Leu-[CF₂-Val]Ile-His-OH,
 5 MeOSuc-Pro-Phe-His-Leu-[CF₂-Val]Ile-His-OH,
 MeOSuc-His-Phe-His-Leu-[CF₂-Val]Ile-His-OH,
 MeOSuc-His-Pro-Phe-His-Leu-[CF₂-Val]Ile-OH,
 MeOSuc-His-Pro-Phe-His-Leu-[CF₂-Val]His-OH,
 BOC-His-Pro-Phe-His-Leu[CF₂-Val]-Ile-His-Oh,
 10 BOC-His-Pro-Phe-His-Leu-[CF₂-CO]-Ile-His-NH₂,
 BOC-Pro-Phe-His-Leu[CF₂-Val]-Ile-His-NH₂.

The compounds of Formula (Io) inhibit renin and
 therefore are used as antihypertensive agents useful in
 treating hypertension. For their end-use application, the
 15 potency and other biochemical parameters of the enzyme
 inhibiting characteristics of the compounds of (Io) are
 readily ascertained by standard biochemical techniques
 well known in the art. Actual dose ranges for their spe-
 cific end-use application will, of course, depend upon the
 20 nature and severity of the disease state of the patient or
 animal to be treated as determined by the attending diag-
 nostician. It is to be expected that the general end-use
 application dose range will be about 0.01 to 10 mg per kg
 per day for an effective therapeutic effect.

25 The compounds of Formula I which are useful as inhi-
 bitors of pepsin are represented by the structural formula



30 with the proviso, that the depicted carbonyl moiety
 attached to X may exist in its chemically reduced form,



wherein X generically is X₁ and X₂, which when X is X₁

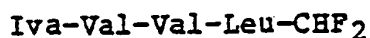
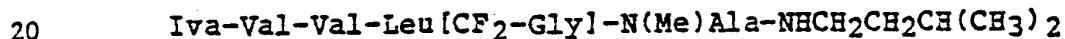
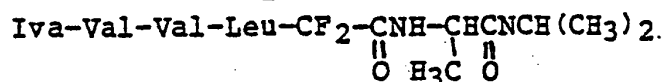
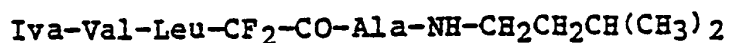
5 -CF₂H, CF₃ and CONH₂ are preferred, when X is X₂
then,

R₄ is selected from the Groups E, G and F with Gly being preferred,

10 R₅ is selected from Groups E and F with Ala being preferred and Y is -NECH₂CH(CH₃)₂ or -NECH₂CH₂CH(CH₃)₂,

R₁ is -P₂-P₃-P₄ with P₂ being selected from Groups E or F with Val being preferred, P₃ is selected from Groups E or F with Val being preferred or is deleted and P₄ is selected from Group K preferably Iva, and

15 R₂ is selected from Groups E and F with Leu being preferred. The preferred compounds are:



The compounds of the formula (Ip) inhibit pepsin and therefore exert an antiulcer effect useful in the treatment and prevention of ulcers. For their end-use application, the potency and other biochemical parameters of the enzyme inhibiting characteristics of the compounds of (Ip) are readily ascertained by standard biochemical techniques well known in the art. Actual dose ranges for their specific end-use application will, of course, depend upon the nature and severity of the disease state of the patient or animal to be treated as determined by the attending diagnostician. It is to be expected that the general end-use

25

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application dose will be about 0.01 to 0.1 mg per kg per day for an effective therapeutic effect.

The compounds of formula I which are useful as inhibitors of Cathepsin D are represented by the structural formula



wherein X generically is X₁ or X₂ which when X is X₁, the preferred groups are -CO₂R₃ or -CF₃, and when X is X₂,

R₄ is selected from Groups E and F with Phe being preferred,

R₅ is selected from Groups E and F with Ala being preferred,

Y is -NH(CH₂)₂CH(CH₃)₂ or -NHCH₂CH(CH₃)₂,

R₁ generically is -P₂-P₃-P₄ with P₂ being selected from Groups E and F, with Val being preferred, P₃ is selected from Groups E and F with Val being preferred, and P₄ is selected from Group K with CBZ being preferred, and

R₂ is selected from Groups E and F, with Phe being preferred. The preferred compounds are:

CBZ-Val-Val-Phe-CF₂-CO-Ala-Iaa

CBZ-Val-Val-Phe-CF₂H,

CBZ-Val-Val-Phe-CF₃,

CBZ-Val-Val-Phe[CF₂-Phe]Ala-NH(CH₂)₂CH(CH₃)₂,

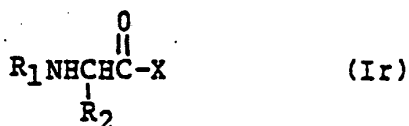
CBZ-Val-Val-Phe[CF₂-Phe]Ala-NHCH₂CH(CH₃)₂,

Iaa being isoamyl amide.

As inhibitors of Cathepsin D the compounds of formula (Iq) are useful for the same end-use applications set forth for human leukocyte elastase inhibitors (Ia) and are also useful as antidemyelinating agents useful to prevent and arrest nerve tissue damage. For their end-use application, the potency and other biochemical parameters of the enzyme inhibiting characteristics of the compounds of (In) are readily ascertained by standard biochemical techniques well known in the art. Actual dose ranges for their specific end-use application will, of course, depend upon the nature and severity of the disease state of the patient or animal to be treated as determined by the attending diagnostician. It is to be expected that the general end-use application dose range will be about 0.01 to 10 mg per kg per day for an effective therapeutic effect.

The compounds of formula I which are useful as inhibitors of angiotension converting enzyme (ACE) are represented by the structural formula

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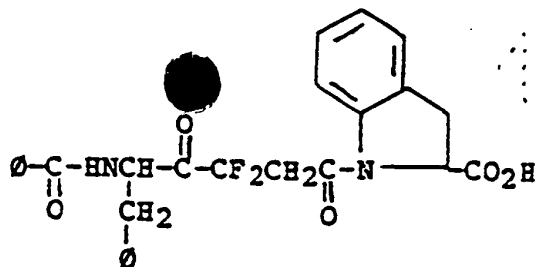
wherein X is only X₂ wherein,

R₄ is selected from Groups E or G with Gly being preferred,

R₅ is selected from Groups A, B, C, D, E, F and G with Group D being preferred and Y is OH,

R₁ is selected from Group K with Bz being preferred,

R₂ is selected from Group E, F and G with Phe being preferred. The preferred species are illustrated as



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5 with \emptyset being phenyl. This preferred compound is also shown as Bz-Phe[CF₂-Gly]Ind-OH.

Other preferred compounds are -

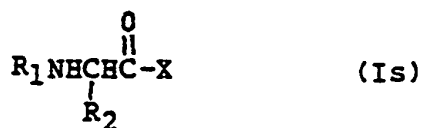
Bz-Phe[CF₂-Gly]Pro-OH

Bz-Phe-CF₂-CO-Pro-OH

10 Bz-Phe-[CF₂-Gly]-Pro-OH

The compounds of formula (Ir) inhibit ACE and are therefore useful as antihypertensives useful in treating hypertension. For their end-use application, the potency and other biochemical parameters of the enzyme inhibiting
15 characteristics of the compounds of (Ir) are readily ascertained by standard biochemical techniques well known in the art. Actual dose ranges for their specific end-use application will, of course, depend upon the nature and severity of the disease state of the patient or animal to
20 be treated as determined by the attending diagnostician. It is to be expected that the general end-use application dose range will be about 0.01 to 10 mg per kg per day for an effective therapeutic effect.

The compounds of Formula I which are useful as inhibitors of enkephalinase are represented by the structural
25 formula

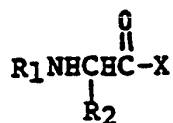


wherein X generically represents X₂ wherein,
R₄ is selected from Group E or F with Phe being preferred,
R₅ is selected from the Groups E or F or zero with the
proviso that when R₅ is zero, Y is NH₂, with Met
being preferred and Y is NH₂ or OH, preferably OH
when R₅ is Met or other α -amino acid,
R₁ generically is -P₂-P₃, with P₂ being Gly and P₃ being
selected from Group F or is deleted with Tyr being
preferred, and
R₂ is Gly. The preferred compounds are:

H-Tyr-Gly-Gly-CF₂-CO-Phe-Leu-OH,
H-Tyr-Gly-Gly[CF₂-Phe]Met-OH
H-Tyr-Gly-Gly[CF₂-Phe]LeuNH₂
H-Tyr-Gly-Gly-CF₂-CO-Leu-OH

The compounds of formula (Is) inhibit enkephalinase
and therefore are useful as analgesics. For their end-use
application, the potency and other biochemical parameters
of the enzyme inhibiting characteristics of the compounds
of (Is) are readily ascertained by standard biochemical
techniques well known in the art. Actual dose ranges for
their specific end-use application will, of course, depend
upon the nature and severity of the disease state of the
patient or animal to be treated as determined by the
attending diagnostician. It is to be expected that the
general end-use application dose range will be about 0.01
to 10 mg per kg per day for an effective therapeutic
effect.

The compounds of Formula I which are useful as inhi-
bitors of Pseudomonas elastase are represented by the
structural formula



(It)

wherein X generically is X₂ with,

5 R₄ being selected from Groups E and F with Ile being preferred,

R₅ is selected from Groups E and G with Ala being preferred and Y is NH₂,

10 R₁ is -P₂-P₃ with P₂ being selected from Group E with Ala being preferred, P₃ is selected from Group K with MeOSuc being preferred,

R₂ is selected from Groups E and G with Ala being preferred. The preferred compounds are:

MeOSuc-Ala-Ala[CF₂-Ile]Ala-NH₂

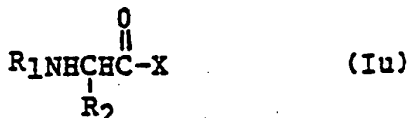
15 MeOSuc-Ala-Ala-CF₂-CO-Ile-Ala-NH₂

The compounds of the Formula (It) inhibit Pseudomonas elastase and therefore are useful as antibacterial agents particularly useful against infections caused by Pseudomonas bacteria. For their end-use application, the potency and other biochemical parameters of the enzyme inhibiting characteristics of the compounds of (It) are readily ascertained by standard biochemical techniques well known in the art. Actual dose ranges for their specific end-use application will, of course, depend upon the nature and severity of the disease state of the patient or animal to be treated as determined by the attending diagnostician. It is to be expected that the general end-use application dose range will be about 0.01 to 10 mg per kg per day for an effective therapeutic effect.

20

25

The compounds of formula I which are useful as inhibitors of leucine aminopeptidase are represented by the formula



wherein X generically embraces all of X₁ and X₂ which when X is X₁, CO₂R₃ or -CF₃ are preferred and when X is X₂,

10 R₄ and R₅ each are any Group except K with Ala and Group E being preferred, Y is NH₂,

R₁ is hydrogen, and

R₂ is selected from Groups A, B, E, F and J with Phe, Leu, Glu and with Arg being preferred. The preferred compounds are:

H-Leu-CHF₂

H-Leu-CF₂-COOEt

H-Arg-CF₃

H-(p-gua)Phe-CF₃

20 H-Leu-CF₃ and H-Leu-COOH,

H-Leu[CF₂-Ala]Ala-NH₂ and H-Leu-COOMe or Leu-COOH

H-Arg-CO-CF₂-Phe-OH

The compounds of formula (Iu) are inhibitors of leucine amino peptidase and therefore are useful as immunostimulants useful in conjunctive therapy in the treatment with other known anticancer agents. For their end-use application, the potency and other biochemical parameters of the enzyme inhibiting characteristics of the compounds of (Iu) are readily ascertained by standard biochemical techniques well known in the art. Actual dose ranges for their specific end-use application will, of course, depend upon the nature and severity of the disease

state of the patient or animal to be treated as determined by the attending diagnostician. It is to be expected that the general end-use application dose range will be about 0.01 to 10 mg per kg per day for an effective therapeutic effect.

The compounds of Formula I which are useful as inhibitors of kallikreins, tissue or plasma, are represented by the formula



wherein X is X₁

R₂ preferably is the Arg residue,

R₁ is a peptide -P₂P₃ with

P₂ being selected from the Groups F and E with Phe being preferred

P₃ being selected from Groups C, E or F, the residues of which may be in either the D- or L-configuration. The preferred compounds of this formula (Iv) are:

D-Pro-Phe-Arg-CF₂H

D-Pro-Phe-Arg-CF₃

D-Pro-Phe-Arg-CO₂H

D-Pro-Phe-Arg-CONH₂

The compounds of formula (Iv) are inhibitors of the kallikreins, tissue or plasma, and therefore inhibit kinin formations. Kinins, generally known to induce pain and vascular permeability associated with inflammation and infection, e.g. bacterial and viral, the inhibition of the kinin formation renders these compounds useful in the

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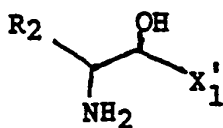
alleviation of pain and inflammation. Furthermore, these compounds are useful as male contraceptives in that they will dramatically interfere with normal sperm function. In their end-use application dose range will be about 0.01
5 to 10 mg per kg per day for an effective therapeutic effect.

Having defined the scope of compounds embraced within the generic formula I and within the individual sub-generic groups of each of the 21 enzymes, the manner in which such compounds may be prepared will hereinbelow be
 5 described as illustrated.

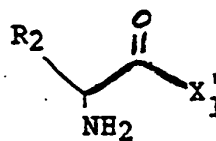
The preparation of the compounds of formula I may be achieved using standard chemical reactions analogously known to be useful for the preparation of a variety of known peptides. Indeed, for the most part, once certain
 10 key intermediate α -amino acid derivatives are prepared, the procedures for obtaining the final products may readily be effected using standard techniques known to those skilled in the field of peptide chemistry. For this purpose, a handy reference text for these techniques is
 15 the 1985 "The Practice of Peptide Synthesis" by M. Bodanszky and A. Bodanszky, wherein the parameters and techniques affecting the selection, use and removal of protective groups for individual and groups of amino acids is detailed, and which also contains activation and
 20 coupling techniques and other special procedures. However, before the application of these peptide chemistry techniques may be applied, certain key intermediates containing the activated electrophilic ketone moiety must first be prepared. The preparation of the key
 25 intermediates is described as follows.

For those compounds wherein X_1 represents either $-CF_2H$ or $-CF_3$, the key intermediates required for the application of the standard peptide coupling techniques are compounds of formula IIIa-b

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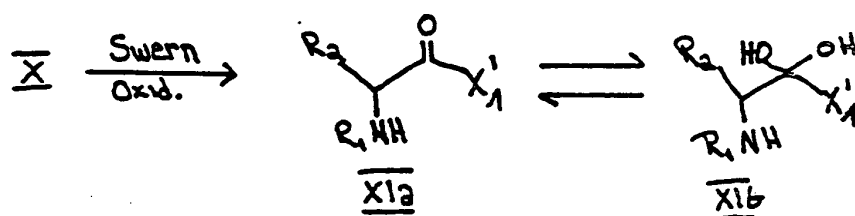
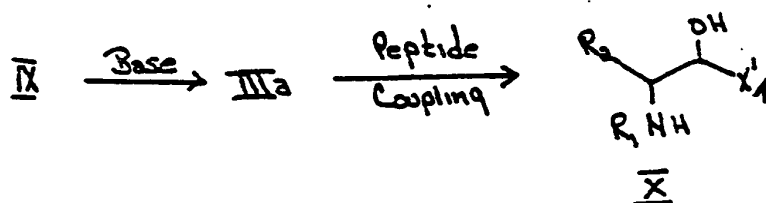
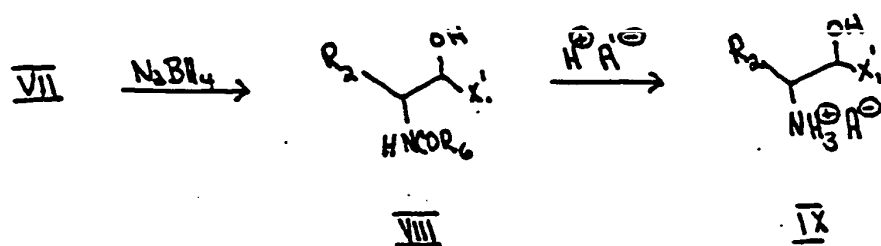
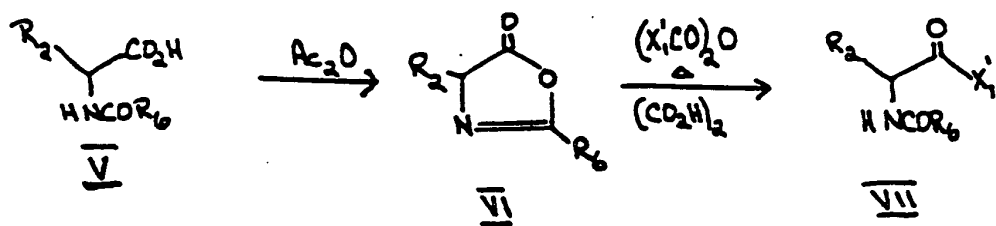
IIIa



IIIb

wherein X_1' is $-CF_3$ or $-CF_2H$, and R_2 is as previously defined in formula I. Similarly, designations R_1 , R_2 , R_3 , R_4 , R_5 and Y shown in the following reaction schemes A through D are as defined in formula I, except that any
5 subgeneric or other modifications thereof (as in X_1') are highlighted by the use of a primed symbol with a specific designation for such modified symbol. The preparation and application of these compounds is depicted by Reaction Scheme A.

REACTION SCHEME A



wherein R_6 is alkyl, phenyl or other equivalent moiety,
and X_1 is $-CF_2H$ or $-CF_3$.

In general, the formation of the substituted azlactones (VI) is effected from the N-protected amino acids (V) by standard reaction conditions wherein the amino acid derivative (V) is heated in the presence of an acid anhydride. The so-produced azlactone (VI) is reacted with a di- or trifluoroacetic acid anhydride or acid halide to give a fluorinated intermediate which (with or without isolation) is treated with anhydrous oxalic acid to produce the N-protected fluorinated ketone (VII) whereupon the ketone is chemically reduced to its alcoholic amide VIII. The amide (VIII) is cleaved under standard acidic conditions to yield its amide acid salt (e.g., its hydrochloride (IX)). After neutralization, the alcohols (IIIa) may be coupled to R_1 using standard peptide chemistry techniques to produce compounds (X) which are subjected to the Swern oxidation procedure to obtain the desired products XIa and XIb (the ketone or hydrate respectively). Alternatively, the alcohols (IIIa) may be oxidized to the ketones (IIIb) which are coupled to R_1 according to standard peptide chemistry techniques. When employing this alternative route, the amino moiety is first protected with a Boc protecting group, the OH function oxidized to its ketone via Swern oxidation procedures, and then the Boc protecting group removed and the resulting compounds (IIIb) are then coupled to R_1 .

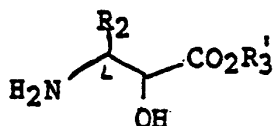
In effecting the foregoing reactions, standard and well-known techniques analogously known are utilized, for example, the azlactones (VI) are chemically transformed to their di- or trifluoromethyl derivatives (their X_1 derivatives) (VII) by heating the azlactone and the fluoroacetic

acid anhydride or acid halide reactants at temperatures of about 30° to 200°C for about 1-24 hours (although under very mild conditions it may take up to one week) preferably using molar equivalent quantities of the reactants. In the event excess quantities of the anhydride reactant are used, such excess should be removed before the next step and the crude product is treated with anhydrous oxalic acid. The fluorinated ketone is reduced to its alcohol using excess quantities of sodium borohydride or any other suitable reducing agent, e.g., sodium cyanoborohydride. Following reduction, the reaction is quenched and the amide is cleaved under standard acidic conditions in water, alcohol or other hydroxylic solvent. The solution is made basic and extracted to obtain the corresponding alcohol (IIIa).

It is, of course, obvious to one of ordinary skill in the art that the conditions of the steps of Reaction Scheme A may have an impact on the R₂ side chain and thus procedures will have to be taken to protect those R₂ moieties which are not compatible with the various reaction conditions. For example, R₂ moieties which belong to Group E are generally compatible. Similarly, R₂ side chain radicals from Groups F, J, and G are compatible. Radicals of Group A need protection. Since arginine may be considered as a derivative of ornithine, the ornithine derivative may first be prepared and then converted to the arginine side chain otherwise the guanidino function of the arginine moiety will have to be protected. The Group C radicals of serine, threonine and cysteine must be protected, preferably with an ether (e.g., benzyl ether). Preferably, the -OH and -SH functions of these groups are protected before the azlactone is formed. The (X) intermediate wherein X₁¹ is -CF₃ and R₂ is H is known (Journal of the American Chemical Society, 70, 143

(1948)), and thus the CF_2H analogs may be prepared using the analogous procedures. The carboxyl moiety of the Group B R_2 side chains must also be protected. The need for the selection of the protecting groups and the
 5 reaction conditions for compatability, selective cleavage and other factors obvious to those skilled in the art are well known and appreciated by those skilled in this field of chemistry.

For those compounds wherein X_1 represents CO_2R_3 ,
 10 CONR_3 or COR_5Y , the key intermediates required for the application of the standard peptide coupling techniques have the formula

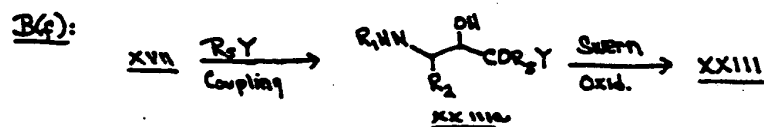
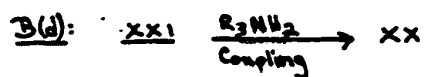
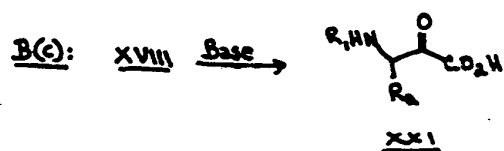
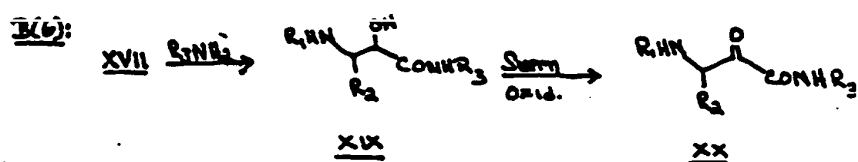
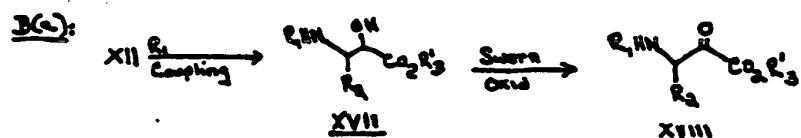
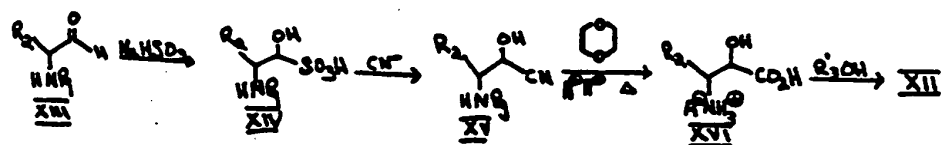


XII

with R_3' being as previously defined for R_3 except that H is omitted from its definition.

20 The preparation and application of these compounds may be depicted by the following reaction scheme. (NB. The desired stereochemistry at the N-substituted carbon is obtained unless otherwise noted.)

REACTION SCHEME B



wherein A^- is the anion of the acid employed and R_3^1 , except for the exclusion of H, is as defined for R_3 , and Pg is a suitable amino moiety protecting group, and in XXV of reaction (b) R^1 is the same as $R_5C(=O)-$, and R^5 is other than zero.

Compounds XIII are generally known starting materials. (In general, such materials may be prepared by converting the appropriate L-amino acid to its ester, preferably its methyl ester by standard esterification procedures (e.g., $SOCl_2$ in the presence of alcohol), the ester is N-protected, preferably with di-t-butyl dicarbonate (Boc). The so-protected R_2 -amino acid is chemically reduced to the desired aldehydic form (XIII) of the N-protected amino acid, preferably with diisobutylaluminum hydride (Dibal); all such procedures being by techniques well known in the art. Of course, other procedures for obtaining the same end products are readily available by techniques well known in the art.

The reaction steps of Reaction Scheme B also use standard and well known techniques. For example, conversion of the aldehydes (XIII) to the corresponding cyanohydrin is effected by reaction with a bisulfite (e.g., $NaHSO_3$) to obtain a bisulfite addition product XIV which is treated with a metalocyanide (e.g., KCN) to produce the corresponding cyanohydrin (XV). Of course, other procedures are readily available for the conversion of the aldehyde to the cyanohydrin derivatives (XV). The cyanohydrins are heated in a mixture of an aqueous acid and water miscible solvent such as an ether (preferably dioxane) to form the desired hydroxy acid (XVI) as a mixture

of its diastereoisomers. These compounds are subjected to neutralization and purification by standard procedures, such as ion exchange resin chromatographic techniques, to yield products which are esterified to obtain the desired key intermediates (XII).

In effecting B(a), the amino esters (XII) are coupled with the R_1 moiety according to standard peptide coupling techniques, taking care that the protective group utilized (if any) are selectively cleavable over the CO_2R_3' moiety. Oxidation of the coupled products is best achieved with the Swern reaction to convert the alcohol to its keto form which, as noted above, can exist in equilibrium with its hydrate.

In effecting B(b), the ester (XVII) is treated with an amine (R_3NH_2) to give the amide (XIX) which may be oxidized via the Swern oxidation procedures to its keto amide (XX). In this instance oxidation conditions other than that of the Swern conditions may also be employed (e.g. oxidation with pyridinium dichromate).

In effecting B(c), the keto ester (XVIII) is treated with a base (e.g., LiOH) to give its keto acid (XXI) except in the case when R_1 contains a terminal methoxysuccinyl moiety in which case the R_3' must be selectively cleavable, e.g., R_3' ought to be t-butyl or benzyl which are selectively cleaved under acidic or hydrogenolysis conditions respectively. In effecting step B(d) the keto acid (XXI) is converted to its amide by standard coupling procedures.

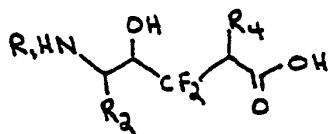
In effecting B(e), the acid is coupled with R_5Y according to standard procedures, taking care to protect and deprotect as needed in the light of the various groups

in the definition of R_5 and Y . In reaction B(f) the ester (XVII) is converted to an amide and the amide oxidized according to Swern oxidation conditions.

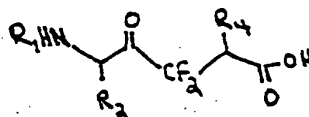
As noted above (following the discussion of Reaction Scheme A), the conditions of the foregoing reactions in the obtention of the desired intermediates and final products of Reaction Scheme B are effected taking care that the R_2 -side chain radicals are compatible. Although Groups E, F and G are generally compatible, protection of some R_2 side chains will be necessary. For example, histidine will have to be protected, while tyrosine and tryptophan ought be protected in order to improve overall yields. Again, ornithine must have its terminal delta amino group protected and ornithine may be converted to arginine. A protecting group would also be needed on a guanidino group. All amino acids, eg. cysteine and threonine having reactive groups in their side chain preferably are protected.

In the preparation of those compounds wherein X_2 is CF_2CHCOR_5Y , the intermediates will be those compounds

$\begin{matrix} R_4 \\ | \\ CF_2CHCOR_5Y \end{matrix}$
of the formulae

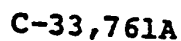


XXIVa



XXIVb

which compounds may be prepared and applied according to Reaction Scheme C.



The azlactone (VI) is prepared as in Reaction Scheme A and the steps going from VI to XXV are analogous to those of that Scheme except that the azlactone is treated with an unsaturated fluorinated carboxylic acid anhydride, and the product is treated with anhydrous oxalic acid to produce compounds XXV. Utilizing these intermediates (XXV) several pathways may be utilized to obtain the desired products. In one pathway (C-1) the intermediate is sequentially subjected to a) a chemical reduction of the electrophilic ketone, and b) hydrolysis of the amide by standard procedures prior to coupling the R₁ moiety. The coupling is readily accomplished using standard peptide chemistry coupling procedures (already referenced) to produce compounds of formula XXVI these intermediates are also available for alternate pathways, i.e., C-1a or C-1b. In pathway C-1a the intermediates are sequentially treated with a) ozone according to standard ozonolysis procedures to produce the corresponding ozonide, b) treatment with dimethylsulfide to convert the ozonide to its aldehydic form and (c) oxidation, preferably using the Jones oxidation procedure to produce compounds of formula XXIVa. These compounds (XXIVa) are sequentially subjected to a) R₅Y coupling and b) Swern oxidation reactions according to already described standard procedures. In those instances wherein it is desired to first protect the hydroxy group, pathway C1-b is available. This pathway essentially mimics C-1a except that the hydroxy function is protected prior to ozonolysis and the hydroxy protecting group (i.e. R₇) is removed in preparation for the Swern oxidation reaction. Typical protecting groups compatible with the described reactions can be used. Preferably the methoxyethoxymethyl group is utilized and is readily removed prior to the Swern oxidation by standard techniques such as by treatment with Zn Br₂.

In pathway C-2 the intermediates are direct
 subjected to the above described sequential reactions
 (ozonolysis, treatment with DMS, and oxidation) to produce
 compounds of formula XXIX which in preparation for R₅Y
 5 coupling are converted to the spiro lactones of formula
 XXX. Coupling, and deprotection (if necessary) produce
 the desired compounds XXXI using standard techniques.

Pathway C-3, a variation of pathway C-1b, first
 reduces the electrophilic ketone to a hydroxy function
 10 which is protected with a suitable protecting group (e.g.
 methoxyethoxymethyl) and the resulting compounds XXVI are
 subjected to (a) ozonolysis, treatment with DMS, and the
 Jones oxidations, (b) R₅Y coupling and deprotection
 reactions, and (c) Swern oxidation, (all of these
 15 reactions being done according to the above described
 procedures for these reactions) to produce the desired
 compounds of formula XXXI.

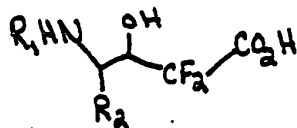
In those instances wherein difluorinated acid anhy-
 drides are required for the preparation of the fluoro-
 20 methyl azlactones, such anhydrides may be prepared by
 reacting tetrafluoroethylene (F₂C:CF₂) with an
 R₄CH=CHCH₂OH reactant in the presence of a base (e.g. NaH)
 and the so-desired R₄CH=CHCH₂OCF₂-CF₂H intermediate is
 treated with butyllithium to produce an acid fluoride
 25 (FC-CF₂CHR₄CH=CH₂) which is converted to its anhydride

$$\begin{array}{c} \text{FC}-\text{CF}_2\text{CHR}_4\text{CH}=\text{CH}_2 \\ \parallel \\ \text{O} \end{array}$$

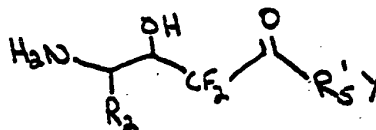
 by standard procedures. Here again, it is obvious that
 compatibility features must be facilitated to ensure that
 the relevant groups can withstand the butyllithium
 30 reaction; thus the R₄ moiety has to be protected when
 incompatible with the butyllithium reaction.

For those compounds wherein X₂ represents $\text{CF}_2\text{C}(\text{R}_5)\text{R}_6$

the key intermediates useful for the preparation of the compounds of formula I bearing this group will be of the formulae



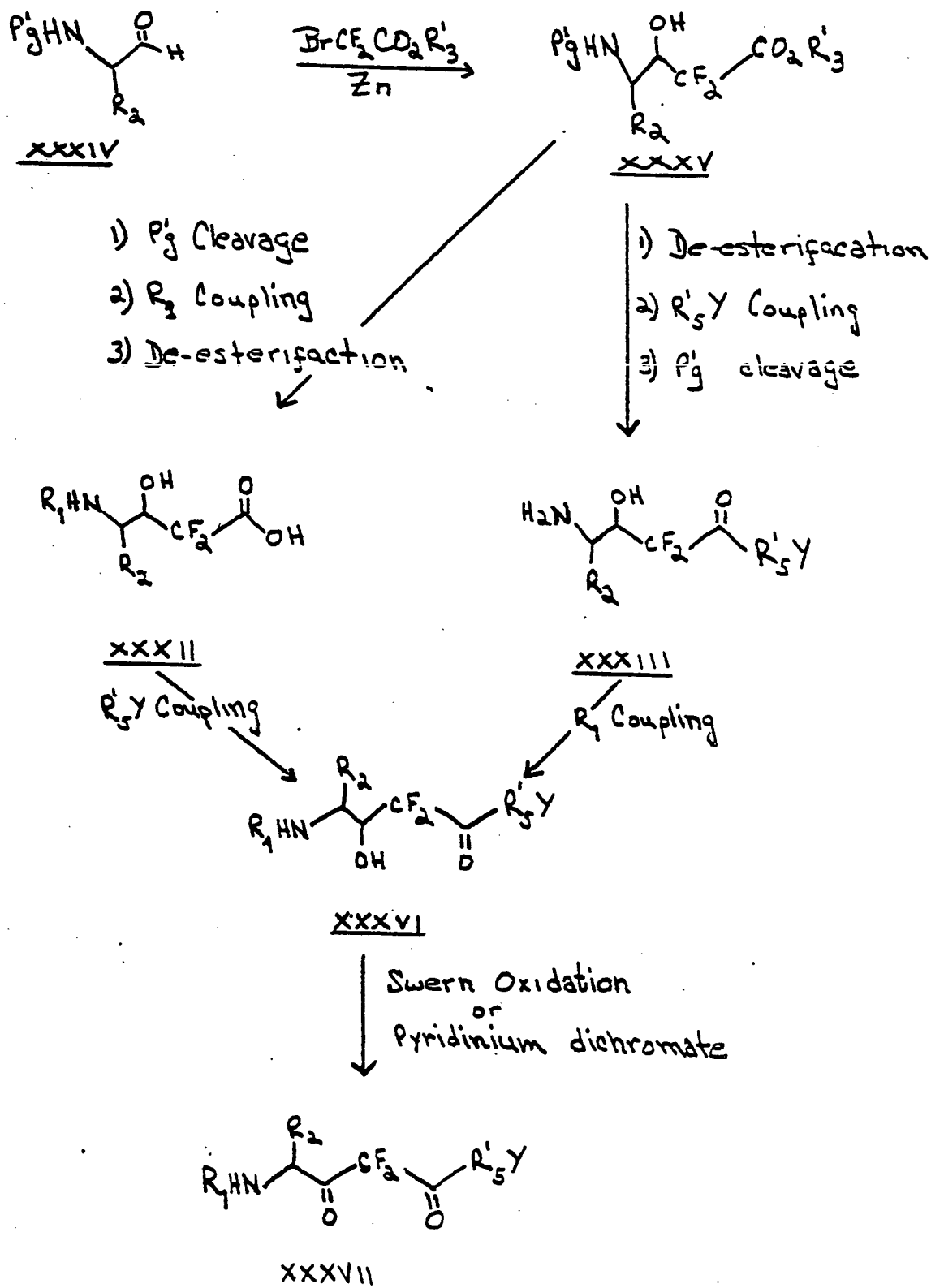
XXXII



XXXIII

the preparation and application of which is depicted by Reaction Scheme D.

REACTION SCHEME D



wherein P_g^1 is a protecting group suitably stable to the reaction steps involved and which is selectively cleaved following R_5Y coupling and the initial alkylation of XXXIV, R_5^1Y is as defined for R_5Y except that R_5 optionally contains a protecting group.

- 5 Following the initial step of alkylating the N-protected aldehyde of formula XXXIV with an appropriate halide in the presence of zinc, the steps following the obtention of compounds XXXV follow the analogous procedures already described for Reaction Schemes A-C.

Having generally described the procedures by which the compounds of this invention may be prepared, the following specific examples serve to further illustrate the standard techniques and procedures which may be utilized to prepare the enzyme inhibitors of this invention (I).

EXAMPLE 1

N1-(2-Hydroxy-3,3-difluoro-1-isobutyl-5-hexenyl)-N2-isovaleryl valinamide

To a solution of 0.621 g (3 mmol) of 6-amino-4,4-difluoro-8-methyl-1-nonen-5-ol (obtained from the corresponding HCl salt by treatment of aqueous solution with 4N NaOH and extracted with Et₂O) and 0.603g (3 mmol) of N-isovaleryl valine in 30 ml THF at 0°C was added 0.62 g of dicyclohexyl carbodiimide. The mixture was stirred for 15 hours at 25°C, filtered and the filtrate concentrated to yield a semisolid which was dissolved in CH₂Cl₂. The organic layer was washed with 1N HCl aqueous KHCO₃ and then brine, dried (MgSO₄) and flash evaporated to afford a solid, which was purified further by chromatography in SiO₂ (CHCl₃/Et₂O (2:1)). The product containing fractions were combined, flash evaporated to give the desired product. R_f 0.15 (CHCl₃/Et₂O) (2:1)).

EXAMPLE 2

3-Phenacetyl-amino-1,1,1-trifluoro-2-propanol

To a mixture of 1.3 g of 3-amino-1,1,1-trifluoro-2-propanol HCl and 1.62 g of triethylamine in 26 ml of THF at 0°C under nitrogen was added dropwise a solution of 1.27 g of phenacetylchloride in 5 ml THF. The reaction mixture was allowed to warm to 25°C and then stirred for 1 hour. The mixture was diluted with CH₂Cl₂ and washed with H₂O, twice with 0.1N HCl, and then brine. After drying (MgSO₄), the solvent was flash evaporated and the residue (1.8 g) recrystallized from CH₂Cl₂ to yield 1.4 g of product; M.P. 96°C.

EXAMPLE 3N¹-(2-Hydroxy-3,3,3-trifluoro-1-isopropyl propyl) N²-phenylmethyloxycarbonyl-prolinamide

To a mixture of 1.1 g of N-phenylmethyloxycarbonyl
5 proline TDO-ester (Cf Hollitzer, Seewald and Steglichm Ang.
Int. Edit. 1976, Vol. 15, 444) and 0.42 g of 3-amino-1,1,1-
trifluoro-4-methyl-2-pentanol hydrochloride in 50 ml of
CH₂Cl₂ was added dropwise 0.40 g of triethylamine. After
stirring for 14 hours at 25°C, aqueous KHSO₄ was added, the
10 layers separated and the organic layer washed with aqueous
KHSO₄, aqueous K₂CO₃, H₂O and then brine (2x). After
drying (MgSO₄), solvents were flash evaporated to yield
0.54 g of a solid residue which was recrystallized in Et₂O
to give 0.24 g of the analytically pure expected amide; mp.
15 99°C.

EXAMPLE 43-tert-Butyloxycarbonylamino-1,1,1-trifluoro-5-methyl-2-hexanol

A mixture of 0.5 g of 3-amino-1,1,1-trifluoro-5-
20 methyl-2-hexanol HCl, 0.5 g of di-tert-butyl dicarbonate
and 0.45 g of K₂CO₃ in 1.6 ml in 1.6 ol of H₂O/dioxane
(1:1) was stirred at room temperature for 14 hours. Workup
in Et₂O/H₂O gave, after washing of the organic layer with
aqueous NaHSO₄, H₂O and brine and drying (MgSO₄) followed
25 by flash evaporation of the solvents 0.55 g of the expected
Boc derivative, R_f 0.44 (Et₂O/pentane (1:1)).

EXAMPLE 53-Phenacetyl-amino-1,1,1-trifluoro-2-propanone

To a solution of 0.84 g of oxalyl chloride in 5 ml of
30 CH₂Cl₂ at -55°C (inside temperature) was added 1.03 g of
dimethylsulfoxide in 1 ml of CH₂Cl₂. After 5 minutes at
-55°C, 1.0 g of 3-phenacetyl-amino-1,1,1-trifluoro-2-pro-
panol in 5 ml CH₂Cl₂ and 0.2 ml of DMSO were added. The

mixture was stirred for 15 minutes at -55°C and then triethylamine was added to adjust the pH to 7.0. The mixture was allowed to warm to 25°C, diluted further with CH₂Cl₂ and washed with H₂O and then 1N HCl. The organic layer was
5 dried (MgSO₄), flash evaporated to yield the amide ketone, which was purified further by chromatography on SiO₂ (CHCl₃/Et₂O (2:1)). R_f 0.29. The product containing fractions were combined and solvent was evaporated under reduced pressure to yield the desired product as a mixture
10 of ketone and hydrate.

EXAMPLE 6

3-tert-Butyloxycarbonylamino-1,1,1-trifluoro-5-methyl-2-hexanone

The named product was prepared by the procedure of
15 example 5 except that 3-tert-butyloxycarbonylamino-1,1,1-trifluoro-5-methyl-2-hexanol is used in place of 3-phenactylamino-1,1,1-trifluoro-2-propanol.

EXAMPLE 7

N¹-(2-Oxo-3,3-difluoro-1-isobutyl-5-hexenyl)-N²-isovaleryl 20 valinamide

To a solution of 0.1 ml of oxalyl chloride in 2.5 ml of CH₂Cl₂ cooled to -55°C was added 0.17 ml of DMSO in 2.5 ml of CH₂Cl₂. After stirring for 5 minutes at -55°C, 0.295 g of N¹-(2-hydroxy-3,3-difluoro-isobutyl-5-hexenyl)-
25 N²-isovaleryl valinamide, in a mixture of 2 ml of CH₂Cl₂ and 0.4 ml of DMSO, were added. The mixture was stirred for 15 minutes at -55°C and then triethylamine (about 0.5 ml) was added to adjust the pH of the solution to 8. The mixture was allowed to warm to room temperature, then
30 diluted with CH₂Cl₂ and washed with H₂O, then with 1N HCl. The organic layer was dried (MgSO₄), then flash evaporated to yield 0.270 g of the expected ketone. R_f 0.3 (CHCl₃/Et₂O (2:1)).

EXAMPLE 85-Benzoylamino-3,3-difluoro-4-oxo-6-phenylhexanoic acid

A solution of (2-benzoylamino-4,4-difluoro-7-phenyl-6-hepten-3-one (1.72 g, 5 mmol) in dichloromethane (200 ml) was treated with O₃ at -78°C for 12 minutes (about 6 mmol O₃). Dimethylsulfide (2 ml, 0.033 mol) was added and the solution allowed to warm to room temperature. After removal of solvents (20 Torr, 30°C and 0.05 Torr, 30°C) a slightly colored oil was obtained, which contained free aldehyde, present in about 70% according to H-NMR (CHO versus 2 CH₂).

A solution of the oil in acetone (7.5 ml) was treated with a Jones-solution (7.5 ml, 1 M CrO₃/H₂SO₄) overnight. The organic layer was separated and the aqueous phase extracted with AcOEt (4 x 10 ml). The combined organic layers were washed with brine, dried (MgSO₄) and flash evaporated to yield 1.7 g of the crude acid.

EXAMPLE 9N¹-(2-Oxo-3,3-Difluoro-1-isobutyl-5-carboxypentyl)-N²-isovaleryl-valinamide

The above-named product was prepared from N¹-(2-oxo-3,3-difluoro-1-isobutyl-5-hexenyl)-N²-isovaleryl valinamide by the procedure of example 8.

EXAMPLE 106,6-Difluoro-2-phenyl-4-phenylmethyl-3-aza-1.9-dioxaspiro-(4,4)non-2-en-8-one

A solution of crude 5-benzoylamino-3,3-difluoro-4-oxo-6-phenyl hexanoic acid (1.37 g, 3.8 mmol) in 10 ml THF was kept under N₂ and cooled to 0°C. Pyridine (0.32 ml, 327 mg, 4.1 mmol) was added slowly. After stirring for 10

minutes at 0°C, the solution was cooled further to -10°C, and oxalyl chloride (0.35 ml, 508 mg, 4 mmol) was added. Gas evolution occurred and the mixture was allowed to warm to 0°C, when a second addition of pyridine (0.32 ml, 327 mg, 4.1 mmol) was added slowly. The mixture was warmed to 40°C over 30 minutes, when gas evolution had ceased. AcOEt (60 ml) and water (5 ml) were added, the phases separated, and the organic layer washed with 0.1 HCl, aqueous NaHCO₃, and water (each 2x5 ml). After drying (MgSO₄) the solvents were removed (20 Torr, 40°C) to yield 1.1 g of crude lactone derivative as a yellow colored oil, which crystallized upon addition of hexane. Recrystallization (AcOEt/hexane, 1:10) afforded 830 mg of pure lactone derivative as colorless needles; mp. 145°C.

15

EXAMPLE 11

N-(5-Benzoylamino-3,3-difluoro-1,4-dioxo-6-phenylhexyl)-S-indoline-2-carboxylic acid, phenylmethyl ester

A mixture of indoline-2-carboxylic acid phenylmethylester hydrochloride (1.16 g, .4 mmol) in Et₂O and H₂O (ea. 5 ml) was treated with Na₂CO₃ (solid) and stirred for 10 min. The organic layer was separated and the aq. layer extracted with Et₂O (2 x 10 ml). The combined organic phases were dried (MgSO₄) and flash evaporated (20 Torr, 30°C, and then 0.05 Torr, 30°C). The only residue (960 mg) was dissolved in chloroform (3 ml).

1 ml (about 320 mg, 1.26 mmol of indolinecarboxylic acid phenylmethyl ester of the above solution was added to 6,6-difluoro-2-phenyl-4-phenylmethyl-3-aza-1,9-dioxaspiro-(4,4)non-2-en-8-one (365 mg, 1.06 mmol) dissolved in 1 ml of chloroform. After stirring 40 hr. at 40°C, solvents were evaporated (20 Torr, 30°C) to give an oily residue, which was purified by chromatography on silica gel (10 g, 230-400 mesh, eluent: pentane/AcOEt (20:3)). The product-containing fractions were combined and solvents removed

under reduced pressure to give 500 mg of the expected peptide as an oil.

EXAMPLE 12

N-(5-Benzoylamino-3,3-difluoro-1,4-dioxo-6-phenylhexyl)-
5 (S)-indoline-2-carboxylic acid

A mixture of N-(5-benzoylamino-3,3-difluoro-1,4-dioxo-6-phenylhexyl)-S-proline phenylmethyl ester (500 mg, 0.84 mmol) and 100 mg Pd/C in i-PrOH (30 ml) was hydrogenated under atmospheric pressure for 12 hours at 25°C. The
10 mixture was filtered and the filtrate flash evaporated to yield 350 mg (82%) of the expected acid as a colorless oil.

EXAMPLE 13

2,2-Difluoro-4-pentenoic acid anhydride

A suspension of silver oxide in water was prepared by
15 adding a solution of NaOH (1.76 g, 0.044 mol) in water (100 ml) to an aqueous solution of silver nitrate (7.14 g, 0.042 mol in 100 ml), decanting the supernatant liquid, washing the residue with water (3 x 100 ml) and adding 100 ml of water. To this vigorously stirred suspension was added a
20 solution of 2,2-difluoro-4-pentenoic acid (5.44 g, 0.04 mol) in water (100 ml). After 10 minutes the mixture was filtered and the filtrate concentrated (20 Torr, 30°C) to afford a solid residue; which was dried over phosphorus pentoxide (0.05 Torr, 50°C, 24 hours) to give 8.4 g (87%)
25 of silver 2,2-difluoro-4-pentenoate; a white amorphous powder. A suspension of 7.3 g (0.03 mol) of the silver salt in 50 ml of dichloromethane was stirred under nitrogen, cooled to 0°C and then 1.9 g (1.3 ml, 0.015 mol) of oxalyl chloride was added slowly. The cooling bath was
30 removed and the reaction mixture allowed to warm up to room temperature. Heating to 40°C for 30 minutes completed the reaction. Cooled again to room temperature, the supernatant liquid was decanted and the residue washed with

dichloromethane (2 x 5 ml). The organic layers were combined and the solvents removed by distillation at atmospheric pressure. The so-obtained oily residue was then purified by distillation to yield 2.85 g of very hydroscopic 2,2-difluoro-4-pentenoic acid anhydride, bp 78-80°C./20 Torr.

EXAMPLE 14

2-Benzoylamino-1-phenyl-4,4-difluoro-6-hepten-3-one

A mixture of 2,2-difluoro-4-pentenoic acid anhydride (2.80 g, 0.011 mol) and 2-phenyl-4-phenylmethyl-5(4H)-oxazolinone (2.60 g, 0.0104 mol) was stirred under nitrogen for 20 hours at 60°C (oil bath temperature) to give a lightly red solution. The reaction mixture was then evaporated (0.05 Torr, 40°C) to afford a highly viscous oil, to which under exclusion of moisture, anhydrous oxalic acid (1.0 g, 0.011 mol) was added and the mixture was heated for 15 minutes (110-120°C, oil bath temperature). After the violent gas evolution had ceased, the oil was allowed to cool to 25°C and then dissolved in a mixture of 40 ml of ethyl acetate and 10 ml of water. The organic layer was separated, washed with aqueous sodium bicarbonate (3x), brine, dried over magnesium sulfate and flash evaporated (20 Torr, 30°C) to afford a red oily residue (2.4 g), which was purified further by flash chromatography on silica gel (50 g, 230-400 mesh, pentane/ethyl acetate (3:1)), R_f 0.6. The product-containing fractions were combined and evaporated to give 2.2 g of a solid, which was recrystallized (ether acetate/pentane) to yield 2.04 g of 2-benzoylamino-1-phenyl-4,4-difluoro-6-hepten-3-one as white needles (59%); mp. 98°C.

EXAMPLE 15

6-Benzoylamino-4,4-difluoro-8-methyl-1-nonen-5-one

The above-named product was prepared from 2-phenyl-4-isobutyl-5-(4H)oxazolinone by the same procedure of the preceding example (yield 73% as oil).

EXAMPLE 16N-(3,3,3-trifluoro-2-oxo-1-(phenylmethyl)propyl)benzamide

2.51 g (0.01 mol) of 2-phenyl-4-phenylmethyl-5(4H)oxa-
zolinone and 2.52 g (0.012 mol) of trifluoroacetic anhy-
5 dride are stirred under N₂ for 24 hours at 35-40°C (oil
bath temperature). After cooling to ambient temperature,
the excess of trifluoroacetic anhydride and the acid formed
are flash evaporated (0.01 Torr, 30-50°C). 1.35 g (0.015
mol) of freshly sublimed anhydrous oxalic acid (0.01 Torr,
10 80-100°C) is added and the mixture heated under stirring to
110°-120°C (oil bath temperature). After gas evolution has
ceased (10-15 minutes) the mixture is allowed to cool to
ambient temperature and stirred for about 1-2 minutes with
a mixture of ethyl acetate and H₂O (10/1). Phases are
15 separated and the organic layer washed with a solution of
NaHCO₃ and then brine (each 3 x 20 ml). Drying (MgSO₄) and
flash evaporation (20 Torr and 0.01 Torr/30°C) affords a
solid which can be crystallized from ethyl acetate/hexane
to yield 2.02 g (63%) of the expected trifluoromethyl
20 ketone:hydrate mixture as a white powder; mp. 163°C.

EXAMPLE 17N-[3,3-Difluoro-2-oxo-1-(phenylmethyl)propyl]benzamide

The above-named product was prepared in 50% yield by
the preceding procedure except that difluoroacetic
25 anhydride was used in placed of trifluoroacetic anhydride;
mp. 136°C.

EXAMPLE 18N-[3,3,3-Trifluoro-2-oxo-(4-nitrophenylmethyl)propyl]-
benzamide

30 The above-named product was prepared in 55% yield by
the preceding procedure (example 16) except that 2-phenyl-
4(4-nitro-phenyl)methyl-5(4H) oxazolinone was used in place

of 2-phenyl-4-phenylmethyl-5-(4H)oxazolinone:hydrate mixture; mp. 175°C.

EXAMPLE 19

5 N-[2-(4-Aminoiminomethyl amino phenyl)-1-trifluoroacetyl ethyl]benzamide, hydrochloride

A suspension of 1.77 g (0.0054 mol) of N-benzoyl-(4-guanidino)phenylalanine in 10 ml (1.438g/0.07 mol) of trifluoroacetic anhydride is stirred at 40°C (oil bath temperature) for 20 hours. The clear solution is flash
10 evaporated (0.01 Torr, 40°C) and treated with anhydrous oxalic acid as described in the synthesis of N-[3,3,3-trifluoro-2-oxo-1-(phenylmethyl)propyl]benzamide, to yield 1.2 g (53%) of the expected trifluoromethyl ketone:hydrate mixture as a white powder; mp. 96°C.

15 EXAMPLE 20

N-[3,3,3-Trifluoro-2-oxo-1-(2-methylethyl)propyl]benzamide

The above-named product was prepared in 23% yield by the procedure of example 16 except that 2-phenyl-4-(2-methylethyl)-5-(4H)oxazolinone was used in place of 2-
20 phenyl-4-phenylmethyl-5-(4H)oxazolinone; mp. 94°C.

EXAMPLE 21

N- 3,3,3-Trifluoro-2-oxo-1[(4-phenylmethyloxycarboxamide)-butyl propyl benzamide

The above-named product (as an oil) was prepared in
25 56% yield by the procedure of example 16 except that 2-phenyl-4(4-phenylmethyloxycarboxamido)butyl-5-(4H)oxazolinone was used in place of 2-phenyl-4-phenylmethyl-5-(4H)oxazolinone.

EXAMPLE 22

30 N-(1-Trifluoroacetyl-3-methyl butyl)benzamide

The above-named product (as an oil) was prepared in 33% yield by the procedure of example 16 except that 2-

phenyl-4-isobutyl-5-(4H)-oxazolinone was used in place of 2-phenyl-4-phenylmethyl-5-(4H)-oxzolinone.

EXAMPLE 23

N-(3,3,3-Trifluoro-2-oxo-propyl)benzamide

5 A solution of 7.57 g hippuric acid and 17.4 ml of tri-fluoroacetic anhydride in 60 ml of anhydrous acetone was stirred at 25°C for 16 hours under N₂ to yield a red precipitate, which is isolated by filtration. Refluxing the red solid in 50 ml of H₂O for 1 hour gave a solution, which
10 was extracted with AcOEt. The organic layer was dried (MgSO₄) and flash evaporated to yield crude product which is recrystallized from benzene to give 4.15 g of analytically pure product; mp. 105°C (decomp).

EXAMPLE 24

3-Benzoylamino-1,1,1-trifluoro-2-propanol

A solution of 10 g (0.263 mol) of NaBH₄ in 100 ml of H₂O was added to 14.8 g (59.4 mmol) of N-(3,3,3-trifluoro-2-oxo-propyl)benzamide in 1000 ml of H₂O. After stirring for 2 hours at 25°C, the solution was acidified with
20 concentrated HCl (pH 1), basified by adding NaOH pellets (pH 10) and extracted with AcOEt (3 x 500 ml). After drying (MgSO₄), the organic layer was flash evaporated to give 11 g of a white solid, which was recrystallized from CHCl₃ to yield 10.0 g (72%) of pure trifluoromethyl
25 alcohol; mp. 156°C.

EXAMPLE 25

6-Benzoylamino-4,4-difluoro-8-methyl-1-nonen-5-ol

The above-named product was prepared from 6-benzoylamino-4,4-difluoro-8-methyl-1-nonen-5-one by example 24
30 except that the alcohol was purified by chromatography on silica gel (eluent EtOAc/hexane (1/5)); mp. 110°C.

EXAMPLE 263-Benzoylamino-1,1,1-trifluoro-4-methyl-2-pentanol

The above-named product was prepared from N-(3,3,3-trifluoro-2-oxo-1-(1-methylethyl)propyl)benzamide by
5 example 24 in 77% yield; mp. 150°C.

EXAMPLE 273-Benzoylamino-1,1,1-trifluoro-5-methyl-2-hexanol

The above-named product was prepared from N-(1-trifluoroacetyl-3-methylbutyl)benzamide by the procedure of
10 example 24 in 80% yield.

EXAMPLE 283-Amino-1,1,1-trifluoro-2-propanol hydrochloride

A mixture of 3 g (12.9 mmol) of 3-benzoylamino-1,1,1-trifluoro-2-propanol in 26 ml of H₂O, 26 ml of concentrated
15 HCl and 26 ml of ethanol was refluxed for 20 hours, then concentrated under reduced pressure. The residue was dissolved in water and extracted with diethyl ether. The aqueous layer was then concentrated to give a solid residue which was recrystallized from isopropanol/diethyl ether to
20 yield 1.37 g of the fluorinated amino alcohol.

EXAMPLE 293-Amino-1,1,1-trifluoro-4-methyl-2-pentanol hydrochloride

The above-named product was prepared from 3-benzoylamino-1,1,1-trifluoro-4-methyl-2-pentanol in 75% yield by
25 the procedure of example 28. R_f 0.37 (AcOEt/Et₃N (20:1)).

EXAMPLE 303-Amino-1,1,1-trifluoro-5-methyl-2-hexanol hydrochloride

The above-named product was prepared from the corresponding 3-benzoylamino derivative. mp. 283°C; R_f 0.78
30 (EtOH/NH₄OH, (70/30)) by the procedure of example 28.

EXAMPLE 316-Amino-4,4-difluoro-8-methyl-1-nonen-5-ol hydrochloride

The above-named product was obtained by the procedure of example 28 in 89% yield from the corresponding 6-benzoylamino derivative.

EXAMPLE 324-N-tert-Butoxycarbonylamino 2,2-difluoro 3-hydroxy 6-methyl heptanoic acid ethyl ester

To a refluxing suspension of 0.196 g of activated zinc wool in anhydrous tetrahydrofuran (3 ml) is added a solution of 0.618 g (3mmol) of ethyl bromodifluoroacetate in anhydrous THF (1 ml). After the reaction starts, a solution of 0.5 g (2.32 mmol) of N-tert-butoxycarbonyl leucinal is added. The mixture is left at gentle reflux for 1 hour. The reaction mixture is then cooled, quenched by addition of ethyl acetate (10 ml), brine and 1M KHSO₄. The organic layer is dried over anhydrous MgSO₄, evaporated and purified by flash chromatography (silica gel, 10% EtOAc/cyclohexane) yield, 0.210 g, R_f 0.65 (EtOAc/cyclohexane, 1:1).

EXAMPLE 334-N-tert-Butoxycarbonylamino 2,2-difluoro 3-hydroxy 6-methyl heptanoic acid

A solution of 0.0285 g (0.68 mmol) of LiOH, H₂O in water (2 ml) is added at 0°C to a mixture of 0.210 g (0.62 mmol) of 4-N-~~tert~~-butoxycarbonylamino 2,2-difluoro 3-hydroxy 6-methyl heptanoic acid, ethyl ester in 1,2-dimethoxyethane (DME) (5 ml). The temperature is allowed to raise slowly to room temperature. The mixture is stirred at room temperature overnight. The mixture is diluted with water (10 ml) washed with diethyl ether (10 ml). The aqueous layer is acidified to about pH 2 with 0.1N HCl, extracted with diethyl ether (2x 10 ml). The

combined organic layers is washed with brine and dried over anhydrous MgSO_4 . Filtration and removal of the solvent in vacuo yields the expected acid which is recrystallized from diethyl ether/pentane.

5

EXAMPLE 344-N-tert-Butoxycarbonylamino 2,2-difluoro 3-hydroxy N-(1-isoamylaminocarbonyl ethyl)-6-methylheptanamide

To a mixture of 0.194 g (1 mmol) of alanine isoamyl amide hydrochloride in methylene chloride (or DMF) (5 ml) is added 0.101 g (1 mmol) of triethylamine at 0°C. 4-N-tert-butoxycarbonylamino-2,2-difluoro-3-hydroxy-6-methyl heptanoic acid (0.311 g) and 1-hydroxybenzotriazole (0.202 g) are added, followed by the addition of DCC (0.206 g, 1 mmole) in methylene chloride (5 ml). The reaction mixture is allowed to warm up slowly to room temperature and stirred for 12 hours at that temperature. Dicyclohexylurea is filtered, and the filtrate evaporated under reduced pressure. The residue is dissolved in ethyl acetate, washed successively with cold 1N HCl, 1N NaOH and dried over anhydrous MgSO_4 . The amide is purified by column chromatography (silica gel, EtOAc/cyclohexane; 1:1, Rf 0.22 (EtOAc/cyclohexane, 1:1)).

EXAMPLE 354-Amino 2,2-difluoro 3-hydroxy N-(isoamylaminocarbonyl-ethyl) 6-methyl-heptanamide hydrochloride

4-N-tert-Butoxycarbonylamino 2,2-difluoro 3-hydroxy N-1-isoamylaminocarbonylethyl)-6-methyl heptanamide (0.457 g) is dissolved in a solution of about 4N HCl in diethylether (5 ml) and stirred at room temperature for 14 hours. After the removal of excess reagent under reduced pressure, the solid residue was triturated with ether to form a solid which is dried, in vacuo for 8 hours. Rf 0.63 (AcOH/butanol/ H_2O ; 2:6:2).

EXAMPLE 36

4-[(2-N-isovaleryl-amino-3-methyl-1-oxobutyl)amino]2,2-difluoro 3-hydroxy-N-(1-isoamylaminocarbonylethyl) 6-methyl heptanamide

5 To a solution of 0.130g (0.33 mmol) of 4-amino 2,2-difluoro 3-hydroxy N-(1-isoamylaminocarbonylethyl) 6-methyl heptanamide HCl in THF (10 ml) is added 0.034 g (0.33 mmol) of N-methyl morpholine at room temperature. After 10 min., the mixture is cooled to 0°C; a solution of 0.103 g (0.50 mmol) of DCC in THF (1 ml) is then added, followed by the addition of 0.100 g (0.50 mmol) of N-isovaleryl valine. 10 Stirring is continued for 15 hours, while the temperature is allowed to rise to room temperature. The precipitate is filtered off and the filtrate rinsed with THF. The solvent 15 is evaporated in vacuo; the residue is purified by chromatography (silica gel, CH₃OH/CHCl₃ 2:98) yielding 0.06 g of the expected amide. Rf: 0.45 (CH₃OH/CHCl₃ 8:92).

EXAMPLE 37

20 4-[(2-N-isovaleryl-amino-3-methyl-1-oxobutyl)amino] 2,2-difluoro-N-(1-isoamylaminocarbonylethyl) 6-methyl 3-oxo heptanamide

A solution of 0.214 g (0.40 mmol) of 4-[(2-N-isovaleryl-amino-3-methyl-1-oxobutyl)amino] 2,2-difluoro 3-hydroxy-N-(1-isoamylaminocarbonylethyl) 6-methyl 25 heptanamide in CH₂Cl₂ (4 ml) is added to a suspension of pyridinium dichromate (0.228 g) and 3 Å molecular sieves (0.336 g), containing 20 µl of glacial acetic acid. Stirring is continued for 15 hours at room temperature. Florisil (0.200 g) is added, stirring continued for 0.25 30 hours and the mixture filtered. Removal of the solvent and chromatography (silica gel, ethyl acetate/acetone 7:3)

afford the expected ketone. Rf: 0.3 (ethyl acetate/chloroform 1:1).

EXAMPLE 38

5 4-N-tert-butoxycarbonylamino 2,2-difluoro 6-methyl 3-oxo heptanoic acid, ethyl ester

The title compound was prepared in 65% yield from the alcohol described in example 32 by the preceding procedure. The ketone was purified by chromatography (silica gel, ethyl acetate/cyclohexane 1:9).

10

EXAMPLE 39

4-amino 2,2-difluoro 6-methyl 3-oxo heptanoic acid, ethyl ester hydrochloride

15 The BOC protecting group of the ketone of example 38 is cleaved using the same procedure as for the amide described in example 35. Mp: 127-128°C (decomp).

EXAMPLE 40

Ethyl-3-keto-2-methyl-4,4,4-trifluorobutanoate

20 Sodium hydride (7.05 g of a 50% oil dispersion, 0.15 mol) was washed 3 times with 25 ml of dimethoxyethane to remove the oil and then suspended in 220 ml of dimethoxyethane, under an argon atmosphere and cooled in an ice bath. A solution of ethyl 3-keto-4,4,4-trifluorobutanoate (25.77 g, 0.14 mol) in 25 ml of dimethoxyethane was added dropwise from an addition funnel to the stirred suspension.

25 After the addition was completed, the cooling bath was removed and the reaction mixture stirred for 30 minutes past the cessation of hydrogen gas evolution. Methyl iodide (43.0 ml, 0.70 mole) was added by syringe and the reaction mixture refluxed overnight. The reaction was

30 cooled to room temperature and poured into a separatory

funnel containing a 1:1:1 mixture of saturated ammonium chloride:brine:water. The layers were separated and the aqueous phase extracted with 100 ml ether. The combined organic phase and ether extract was washed with brine,
5 dried over magnesium sulfate and filtered. The solvents were removed by distillation at atmospheric pressure using a Vigreux column leaving ethyl 3-keto-2-methyl-4,4,4-trifluorobutanoate as the pot residue.

Rf: (EtAc/hexane - 20:80)

10

EXAMPLE 41

Ethyl 3-hydroxy-2-methyl-4,4,4-trifluorobutanoate

To a solution of ethyl 3-keto-2-methyl-4,4,4-trifluorobutanoate (20.1 g, 0.10 moles) in 250 ml of absolute ethanol, cooled in an ice bath, was added sodium borohy-
15 dride (1.0 g, 0.25 moles) in portions with stirring. The cooling bath was removed and the reaction stirred at room temperature for 30 minutes. Acetone (5 ml) was added to quench any remaining sodium borohydride and the solvents removed by distillation at atmospheric pressure using a
20 Vigreux column. The residue was diluted with 200 ml of

methylene chloride and poured into a separatory funnel containing 75 ml of a 1:1:1 mixture of saturated ammonium chloride:brine:water. The layers were separated and the aqueous phase extracted with methylene chloride (3 x 25 ml). The combined organic phase and methylene chloride extracts were dried over magnesium sulfate, filtered and distilled at atmospheric pressure to remove the solvents. The residue was then distilled at reduced pressure (20 mmHg) using a Vigreux column to give ethyl 3-hydroxy-2-methyl-4,4,4-trifluorobutanoate, bp 78-84°C, 20mmHg.

EXAMPLE 42

3-Hydroxy-2-methyl-4,4,4-trifluorobutanamide

Into a solution of ethyl 3-hydroxy-2-methyl-4,4,4-trifluorobutanoate (11.4 g, 51.0 mmol) in 85 ml of methanol, cooled in an ice bath, was bubbled in anhydrous ammonia for several minutes. The reaction flask was sealed with a septum and stirred at room temperature for 6 days. The mixture was concentrated using a rotary evaporator and the residue distilled at reduced pressure using a vacuum pump to remove all components with a boiling point less than 25°C at 0.05 mmHg leaving trifluorobutanamide (5.8g, 33.9 mmol) as the pot residue.

EXAMPLE 43

3-Hydroxy-4,4,4-trifluoro-2-butylamine hydrochloride

To potassium hydroxide pellets (15.4 g of 85% pellets, 0.23 moles) dissolved in 45 ml of water and cooled in an ice bath was added bromine (2.7 ml, 51.4 mmol). After several minutes, a solution of 3-hydroxy-2-methyl-4,4,4-trifluorobutanamide (8.0 g, 46.8 mmol) in 45 ml water, pre-cooled in an ice bath, was added. The reaction was stirred at ice bath temperatures for 20 minutes and then at room temperature for 30 minutes. Finally, the reaction was heated on a steam bath for 30 minutes. The reaction

mixture was cooled to room temperature and poured into a separatory funnel where it was extracted with methylene chloride (3 x 50 ml). The aqueous layer was then saturated with solid sodium chloride and extracted further with two portions of methylene chloride (25 ml). The combined organic extracts were dried over magnesium sulfate, filtered and the solvent removed at ambient temperature on a rotary evaporator. The residue was dissolved in anhydrous ether (250 ml) and anhydrous hydrogen chloride gas bubbled through the reaction mixture. A white precipitate formed and the suspension was cooled in an ice bath. The precipitate was filtered and then recrystallized from acetone-ether to give 3-hydroxy-4,4,4-trifluoro-2-butylamine hydrochloride, Rf: 0.25 (NH₄OH/CH₃OH/CH₂Cl₂ - 2:10:88).

15

EXAMPLE 44L-Alanine methyl ester hydrochloride

To a suspension of L-alanine (25.0 g, 0.28 moles) in 125 ml of methanol, cooled in an ice-methanol bath, was added thionyl chloride (21.0 ml, 0.29 moles) dropwise at a rate such that the internal reaction temperature was maintained at 5°C or less. After the addition was completed, the cooling bath was removed and the reaction mixture warmed to 45°C for 2 hours. The reaction mixture was filtered to remove a small amount of yellow solid and the filtrate concentrated using a rotary evaporator. To the resultant oil was added tetrahydrofuran (50 ml) and the mixture evaporated to dryness on a rotary evaporator. The residue was placed under high vacuum to yield an off white solid. Ether (300 ml) was added to the solid and the suspension digested on a steam bath. Cooling and filtering gave L-alanine methyl ester hydrochloride (37.2 g, 0.26 mmol).

EXAMPLE 45Boc-L-Alanine methyl ester

To a stirred suspension of L-alanine methyl ester hydrochloride (10.0 g, 71.6 mmol) in methylene chloride (220 ml) under an argon atmosphere was added triethylamine (10.0 ml, 71.6 mmol). Fifteen minutes later, a solution of di-tert-butyl dicarbonate (15.3 g, 70.2 mmol) in methylene chloride (30 ml) was added dropwise. After the addition was complete, the reaction mixture was stirred at room temperature overnight. The reaction mixture was poured into a separatory funnel containing 50 ml water and the layers separated. The organic layer was washed with 0.5 N hydrochloric acid (2 x 50 ml) and brine (50 ml). The organic layer was then dried over magnesium sulfate, filtered and the majority of solvent evaporated using a rotary evaporator. The last traces of solvent were removed under high vacuum to give Boc-L-alanine methyl ester (14.27 g, 70.2 mmol).

EXAMPLE 46Boc-L-Alaninal

Boc-L-alanine methyl ester (5.0 g, 24.6 mmol) was dissolved in dry toluene (150 ml) under an argon atmosphere and cooled in a dry ice-acetone bath. To this vigorously stirred solution was added a solution of diisobutylaluminum hydride (1.0 M in hexanes, 61.5 ml, 61.5 mmol), pre-cooled in a dry ice-acetone bath, via a transfer needle. After 6 minutes, the reaction was carefully quenched with methanol (4 ml) and the mixture allowed to warm to room temperature. The reaction mixture was poured into a separatory funnel containing 250 ml ether and 200 ml of ice cold 0.25 N hydrochloric acid. The layers were separated and the organic layer was washed with 0.25 N hydrochloric acid (3 x 80 ml) and brine (50 ml). The organic layer was dried over magnesium sulfate, filtered and concentrated using a rotary

evaporator at ambient temperature. The residue was chromatographed using hexanes - 30% ethylacetate to give Boc-L-alaninal. The compound may also be prepared by the technique described in Synthesis, 1983, pg. 676.

5

EXAMPLE 47(3S)-3-Amino-2-hydroxybutanoic acid

To a suspension of Boc-L-alaninal (2.5 g, 14.4 mmol) in ice cold water (30 ml) was added an ice cold solution of sodium bisulfite (1.5 g, 14.4 mmol) in water (10 ml). The resultant suspension was stirred at ice bath temperature overnight. To the resultant solution was added ethyl acetate (200 ml) and then a solution of potassium cyanide (0.9 g, 14.4 mmol) in water (10 ml). The reaction mixture was stirred at room temperature for 4 hours and then poured into a separatory funnel and the layers separated. The organic layer was washed with water (2 x 100 ml) and brine (75 ml), then dried over magnesium sulfate, filtered and concentrated using a rotary evaporator. The cyanohydrin was dissolved in dioxane (50 ml) and concentrated hydrochloric acid (50 ml) added. The reaction mixture was refluxed overnight and then cooled to room temperature. The reaction mixture was poured into a separatory funnel containing ether (100 ml) and the layers separated. The aqueous layer was extracted with a further 100 ml ether and then evaporated to dryness on a rotary evaporator. The resultant residue was dissolved in water (30 ml) and the pH adjusted to approximately 7 with 2N ammonium hydroxide. This solution was placed on a Biorad AG 50 W-X8 H₂O resin column and eluted with 2 N ammonium hydroxide. Combination of the appropriate fractions and evaporation gave crude (3S)-3-amino-2-hydroxybutanoic acid which was then recrystallized from water-acetone to give the desired product (1.05 g, 8.8 mmol) as a white solid.

EXAMPLE 48

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Methyl (3S)-3-amino-2-hydroxybutanoate

5 Into a suspension of (3S)-3-amino-2-hydroxybutanoic acid (1.0 g, 8.4 mmol) in methanol (25 ml) was bubbled anhydrous hydrogen chloride gas until a solution resulted. After a solution resulted, the reaction was cooled in an ice bath and saturated with hydrogen chloride. The cooling bath was removed and the reaction stirred at room temperature for 3.5 hours. The solvent was removed on a rotary
10 evaporator at ambient temperature and the resultant residue dissolved in methanol (25 ml), cooled in an ice bath and saturated with hydrogen chloride gas. Warming of the reaction solution to room temperature and removal of the solvent on a rotary evaporator gave an oil. To this oil
15 was added triethylamine (15 ml) followed by the minimum amount of methanol (about 15 ml) needed to dissolve the initial gummy solid. The solution was cooled in an ice bath and ether (75 ml) added in portions with stirring. The precipitated triethylamine hydrochloride was filtered
20 and the filtrate evaporated to give methyl (3S)-3-amino-2-hydroxybutanoate.

EXAMPLE 49

Boc-L-alanyl-L-proline benzyl ester

25 Boc-L-alanine (19.5 g, 0.10 mol) was dissolved in dry tetrahydrofuran (90 ml) under an argon atmosphere in a flask fitted with an overhead stirrer and an internal thermometer. The solution was cooled to -15°C and N-methylmorpholine (11.4 ml, 0.10 mol) was added followed by isobutylchloroformate (13.4 ml, 0.10 mol) at such a rate as
30 to maintain the internal reaction temperature at -10° to -15°C. Five minutes after the addition was completed, a solution of L-proline benzyl ester hydrochloride (25.2 g, 0.10 mol) and N-methylmorpholine (11.4 ml, 0.10 mol) in chloroform (90 ml) was added dropwise at such a rate as to

maintain the internal reaction temperature at -10° to -15°C . After the addition was completed, the reaction mixture was allowed to slowly warm to room temperature and then stirred at room temperature overnight. The reaction mixture was concentrated using a rotary evaporator, the residue diluted with ethyl acetate (500 ml)/0.2N hydrochloric acid (100 ml) and poured into a separatory funnel. The layers were separated and the aqueous phase extracted with a further 150 ml ethyl acetate. The combined organics were washed with 0.2 N hydrochloric acid (2 x 100 ml), water (100 ml), saturated sodium bicarbonate (2 x 100 ml), again with water (100 ml) and finally brine (100 ml). The organic phase was dried over magnesium sulfate, filtered and concentrated on a rotary evaporator. Chromatography using ethyl acetate-hexane as the eluent gave Boc-L-alanyl-L-proline benzyl ester, R_f : 0.15 (EtOAc/hexane - 30:70).

EXAMPLE 50

L-Alanyl-L-proline benzyl ester hydrochloride

Into a solution of Boc-L-alanyl-L-proline benzyl ester (31.6 g, 83.94 mmol) in ethyl acetate (400 ml) cooled in an ice bath was bubbled hydrogen chloride gas for 15 minutes. The addition of gas was ceased, the cooling bath removed and the solution stirred at room temperature for 1.5 hours. Concentration using a rotary evaporator followed by drying the residue [in a vacuum desiccator over potassium hydroxide pellets overnight] gave L-alanyl-L-proline benzyl ester hydrochloride (25.5 g, 81.5 mmol).

EXAMPLE 51

Boc-L-alanyl-L-alanyl-L-proline benzyl ester

L-alanyl-L-proline benzyl ester hydrochloride (13.0 g, 41.6 mmol) was dissolved in methylene chloride (650 ml) under an argon atmosphere in a flask fitted with an over-

head stirrer. N-methylmorpholine (4.8 ml, 43.6 mmol) was syringed into the solution and, after 5 minutes, Boc-L-alanine (7.9 g, 41.6 mmol) was added followed by 1-ethoxycarbonyl-2-ethoxy-1,2-dihydroquinoline (11.8 g, 47.8 mmol).
5 The resulting solution was stirred at room temperature overnight. The reaction mixture was poured into a separatory funnel containing water (300 ml) and the layers separated. The aqueous layer was extracted with chloroform (200 ml) and the combined organic extracts were washed with
10 0.5 N hydrochloric acid (3 x 200 ml), water (2 x 200 ml), saturated sodium bicarbonate (2 x 200 ml) and brine (200 ml). The organic layer was then dried over magnesium sulfate, filtered and concentrated on a rotary evaporator. Addition of ether-hexane gave crude Boc-L-alanyl-L-alanyl-L-proline benzyl ester which could be recrystallized from
15 ethyl acetate-hexane to give the pure product (15.1 g) mp 124-126°C.

EXAMPLE 52

L-Alanyl-L-alanyl-L-proline benzyl ester hydrochloride

20 Into a solution of Boc-L-alanyl-L-alanyl-L-proline benzyl ester (25.5 g, 57.0 mmol) in ethyl acetate (650 ml) cooled in an ice bath was bubbled hydrogen chloride gas for 15 minutes at which time bubbling was ceased. The cooling bath was removed and the solution stirred at room temperature for 1.5 hours. The reaction mixture was concentrated
25 on a rotary evaporator and the resultant gummy solid dissolved in methylene chloride-hexane. Removal of the solvents gave L-alanyl-L-alanyl-L-proline benzyl ester hydrochloride which was dried over potassium hydroxide
30 pellets in a vacuum desiccator overnight to yield 21.09 (54.7 mmol) of the desired product.

EXAMPLE 53Methoxysuccinyl-L-alanyl-L-alanyl-L-proline benzyl ester

To a solution of L-alanyl-L-alanyl-L-proline benzyl ester hydrochloride (19.2 g, 50.0 mmol), mono-methyl succinate (6.6 g, 50.0 mmol) and 1-hydroxybenzotriazole. \cdot xH₂O (16.9 g) in N,N-dimethylformamide (125 ml) under an argon atmosphere and cooled in an ice bath was added N-methylmorpholine (5.5 ml, 50.0 mmol). After 5 minutes, N,N'-dicyclohexylcarbodiimide (11.9 g, 57.5 mmol) was added, the cooling bath removed and the reaction mixture stirred at room temperature overnight. The reaction mixture was filtered and the filtrate poured into a separatory funnel containing chloroform (750 ml)/0.5N hydrochloric acid (250 ml). The layers were separated and the aqueous phase extracted with chloroform (200 ml). The combined organic extracts were washed with 0.5 N hydrochloric acid (2 x 250 ml), water (2 x 250 ml), saturated sodium bicarbonate (2 x 250 ml) and brine (250 ml). The organic layer was dried over magnesium sulfate, filtered and concentrated on a rotary evaporator. The last traces of solvent were removed using a vacuum pump and the residue chromatographed using acetone-ethyl acetate as the eluent. The resultant crude MeOSuc-L-alanyl-L-alanyl-L-proline benzyl ester was recrystallized from ethyl acetate to give pure product mp 124-127°C.

EXAMPLE 54MeOSuc-L-alanyl-L-alanyl-L-proline

Into a Parr flask flushed with argon containing 4.0 g of 10% palladium on charcoal catalyst was added tert-butanol (775 ml) and then MeOSuc-L-alanyl-L-alanyl-L-proline benzyl ester (13.0 g, 28.2 mmols). The mixture was shaken under 30 psi of hydrogen at 30-40°C overnight. The mixture was filtered through celite and the filtrate concentrated on a rotary evaporator. The residue, was

azeotroped with chloroform-hexane to remove the last traces of tert-butanol and then dried under high vacuum to give MeOSuc-L-alanyl-L-alanyl-L-proline (10.4 g, 28.0 mmol).

EXAMPLE 55

5 Acetyl-L-prolyl-L-alanyl-L-proline benzyl ester

Ac-L-proline (3.05 g, 19.41 mmol) was dissolved in dry tetrahydrofuran (100 ml) under an argon atmosphere in a flask fitted with an overhead stirrer and an internal thermometer. The solution was cooled to -15°C and N-methylmorpholine (2.13 ml, 19.41 mmol) was added followed by isobutylchloroformate (2.51 ml, 19.41 mmol) at such a rate as to maintain the internal reaction temperature at -10° to -15°C. Five minutes after the addition was completed, a solution of L-alanyl-L-proline benzyl ester hydrochloride (6.08 g, 19.41 mmol) in chloroform (70 ml) was added followed by a second portion of N-methylmorpholine (2.13 ml, 19.41 mmol) at such a rate as to maintain the internal reaction temperature at -10° to -15°C. The reaction mixture was then allowed to slowly warm to room temperature and stirred at room temperature for 2.5 hours. The reaction mixture was concentrated to a small residue on a rotary evaporator, diluted with chloroform (500 ml) and poured into a separatory funnel where it was washed with 0.2 N hydrochloric acid (3 x 200 ml) and 5% sodium bicarbonate (200 ml). The organic layer was dried over magnesium sulfate, filtered, concentrated on a rotary evaporator and chromatographed using methylene chloride-methanol as an eluent to give Ac-L-prolyl-L-alanyl-L-proline benzyl ester; Rf: 0.35 (CH₃OH/CH₂Cl₂ - 7:93).

30

EXAMPLE 56

Acetyl-L-prolyl-L-alanyl-L-proline

Into a Parr flask flushed with argon containing 450 mg of 10% palladium on charcoal catalyst was added a solution of Ac-L-prolyl-L-alanyl-L-proline benzyl ester in (1.0 g,

2.41 mmol) absolute ethanol (100 ml). The contents were shaken under 40 psi of hydrogen overnight at room temperature. The mixture was filtered through celite and the filtrate concentrated on a rotary evaporator to give crude Ac-L-prolyl-L-alanyl-L-proline which was crystallized from tetrahydrofuran-methanol-ether to give the desired product (0.57 g) in 73% yield.

EXAMPLE 57

1,1,1-Trifluoro-3-[(N-acetylprolyl)alanylprolylamino]butan-2-ol

Ac-L-prolyl-L-alanyl-L-proline (1.17 g, 3.61 mmol) was suspended in dry acetonitrile (55 ml) under an argon atmosphere in a flask fitted with an overhead stirrer and internal thermometer. The suspension was cooled to -15°C and N-methylmorpholine (0.40 ml, 3.61 mmol) was added followed by isobutylchloroformate (0.47 ml, 3.61 mmol) at such a rate as to maintain the internal reaction temperature at -10° to -15°C . Ten minutes after the addition was completed, a mixture of 3-hydroxy-4,4,4-trifluoro-2-butylamine hydrochloride (0.65 g, 3.61 mmol) and N-methylmorpholine (0.40 ml, 3.61 mmol) in chloroform (25 ml) was added at such a rate as to maintain the internal reaction temperature at -10° to -15°C . The reaction mixture was allowed to warm slowly to room temperature and then stirred at room temperature for 4 hours. The reaction mixture was evaporated on a rotary evaporator to an oily residue which was dissolved in water (65 ml) and treated with a mixed bed resin (J. T. Baker - ANGMI-615, 17 g). After 15 minutes the mixture was filtered and the filtrate evaporated on a rotary evaporator. Chromatography using methylene chloride - 10% methanol as the eluent gave the above-named product, (0.37 g) in 23% yield.

EXAMPLE 581,1,1-Trifluoro-3-[(N-acetylpropyl)alanylpropylaminol]-
butane-2,2-diol

To a solution of oxalyl chloride (72 1, 0.83 mmol) in
5 methylene chloride (1 ml) under an argon atmosphere and
cooled in a dry ice-acetone bath was added dimethylsulfox-
ide (0.12 ml, 1.65 mmol) with stirring. After 5 minutes, a
solution of the compound of example 57 (0.25 g, 0.55 mmol)
in methylene chloride (1.5 ml) was added. The reaction
10 mixture was stirred for 15 minutes, triethylamine (0.50 ml,
3.58 mmol) then added and the reaction warmed to room
temperature for 30 minutes. The reaction mixture was
placed directly onto a silica gel column and eluted with
methylene chloride-methanol. Trituration of the resultant
15 oily solid with ether-hexane and filtration gave the above-
named product, (50 mg, 0.11 mmol).

EXAMPLE 593-[(N-acetylpropyl)alanylpropylaminol]-2-hydroxybutanoic
acid, methyl ester

20 Ac-L-propyl-L-alanyl-L-proline (0.65 g, 2.00 mmol) was
suspended in dry acetonitrile (20 ml) under an argon
atmosphere in a flask fitted with an overhead stirrer and
internal thermometer. The suspension was cooled to -15°C
and N-methylmorpholine (0.22 ml, 2.00 mmol) was added
25 followed by isobutylchloroformate (0.26 ml, 2.00 mmol) at
such a rate as to maintain the internal reaction tempera-
ture at -10° to -15°C. After 10 minutes, a solution of
methyl (3S)-3-amino-2-hydroxybutanoate (0.53 g, 4.00 mmol)
in chloroform (2.5 ml) was added at such a rate as to
30 maintain the internal reaction temperature at -10° to
-15°C. The reaction mixture was slowly warmed to room
temperature and then stirred for 3 hours. The reaction
mixture was evaporated on a rotary evaporator, the residue
dissolved in water (20 ml) and treated with a mixed bed

resin (J. T. Baker - ANGMI-615, 11.0 g). After 15 minutes the mixture was filtered and the filtrate evaporated on a rotary evaporator. Chromatography using methylene chloride-methanol as the eluent gave the above-named product (0.32 g) in 36% yield.

EXAMPLE 60

3-[(N-acetylpropyl)alanylpropylaminol-2,2-dihydroxybutanoic acid, methyl ester

To a stirred solution of oxalyl chloride (0.12 ml, 1.43 mmol) in methylene chloride (1.5 ml) under an argon atmosphere and cooled in a dry ice-acetone bath was added dropwise a solution of dimethylsulfoxide (0.20 ml, 2.86 mmol) in methylene chloride (1.5 ml). After 5 minutes, a solution of the product of example 59 (0.32 g, 0.72 mmol) in methylene chloride (1.5 ml) was added and the reaction mixture stirred for 25 minutes. Triethylamine (0.50 ml, 3.58 mmol) was added and the reaction mixture warmed to room temperature. The reaction mixture was placed directly onto a silica gel column and eluted with methylene chloride-methanol. Evaporation of the appropriate fractions on a rotary evaporator, addition of water (3 ml) to the residue and evaporation gave the above-named product.

EXAMPLE 61

(N-acetylpropyl)alanylpropylaminol-2,2-dihydroxybutanoic acid

To a solution of the product of example 60 (0.10 g, 0.23 mmol) in water (4 ml) cooled in an ice bath was added 1N lithium hydroxide (0.50 ml of an aqueous solution, 0.50 mmol). After 1 hour the pH of the reaction mixture was adjusted to 4.5 to 5.0 with 1N hydrochloric acid and the reaction evaporated on a rotary evaporator. The residue was chromatographed using methylene chloride-methanol as

the eluent. Evaporation of the appropriate fractions, addition of water (2 ml) to the residue and evaporation gave the above-named product (65 mg) in 64% yield.

EXAMPLE 62

5 Boc-L-alanyl-L-alanyl-L-proline

A Parr flask was flushed with argon and charged with 10% palladium on charcoal (0.74 g), followed by the addition of Boc-L-alanyl-L-alanyl-L-proline benzyl ester (1.8 g, 4.0 mmol) dissolved in tert-butanol (300 ml). The
10 reaction mixture shaken under 30 psi of hydrogen at 35°C for 5 hr. After cooling to room temperature, ethanol (50 ml) was added and the solution was filtered through celite and the filtrate concentrated on a rotary evaporator. The
15 residue was azeotroped with chloroform-hexane to remove the last traces of tert-butanol and then dried under high vacuum to give Boc-L-alanyl-L-alanyl-L-proline (1.40 g, 3.9 mmol) in 98% yield.

EXAMPLE 63

20 1,1,1-Trifluoro-3-[(N-tert-butylloxycarbonylalanyl)alanylprolylaminol-4-methylpentan-2-ol]

To a solution of Boc-L-alanyl-L-alanyl-L-proline (1.0 g, 2.80 mmol) in dry acetonitrile (25 ml) was added N-methylmorpholine (0.34 ml, 3.06 mmol). The solution was
25 cooled to -20°C and isobutylchloroformate (0.37 ml, 2.88) was added dropwise. To this solution, a pre-cooled (-20°C) mixture of 3-amino-1,1,1-trifluoro-4-methyl-2-pentanol hydrochloride (0.61 g, 2.91 mmol), N,N-dimethylformamide (4 ml) and N-methylmorpholine (0.34, 3.06 mmol) was added.
30 The reaction mixture was stirred at -20°C for 4 hr, allowed

to warm to room temperature and stirred overnight. Removal of the solvents in vacuo produced a pale yellow residue, which was purified by flash chromatography using ethyl acetate as an eluent to give 1,1,1-Trifluoro-3-[(N-tert-butylloxycarbonylalanyl)alanylproylamino]-4-methylpentan-2-ol (1.09 g, 2.1 mmol) in 76% yield.

EXAMPLE 64

1,1,1-trifluoro-3-[(N-tertbutyloxycarbonylalanyl)alanylproylamino]-4-methylpentan-2-one (180 mg, 0.37 mmol) in 70% yield

A solution of oxalyl chloride (0.078 ml, 0.9 mmol) in methylene chloride (2 ml) was cooled to -55°C and dimethylsulfoxide (0.125 ml, 1.8 mmol) was added dropwise. The solution was stirred for 5 min, followed by the addition of 1,1,1-trifluoro-3-[(N-tert-butylloxycarbonylalanyl)alanyl-proylamino]-4-methylpentan-2-ol (260 mg, 0.53 mmol) in methylene chloride (1.5 ml). The mixture was stirred for 15 min and triethylamine (0.45 ml, 3.2 mmol) was added. The reaction mixture was allowed to warm to room temperature, the solvent was removed in vacuo and the crude product was loaded directly onto a silica gel column (230-400 mesh) for purification. Elution with ethyl acetate gave 1,1,1-trifluoro-3-[(N-tert-butylloxycarbonylalanyl)alanylproylamino]-4-methylpentan-2-one.

EXAMPLE 65

1,1,1-Trifluoro-3-[alanyl-alanylproylamino]-4-methylpentan-2-one

A solution of 1,1,1-Trifluoro-3-[(N-tert-butylloxycarbonylalanyl)alanyl-proylamino]-4-methylpentan-2-one (180 mg, 0.35 mmol) in ethyl acetate (50 ml) was

cooled to 0°C and treated with hydrogen chloride gas for 5 min. The reaction mixture was stirred at 0°C for 1.5 hr, followed by removal of solvent in vacuo. 1,1,1-Trifluoro-3-[alanylalanylprolylamino]-4-methylpentan-2-one (151 mg, 0.34 mmol) was obtained in 96% yield and was used for subsequent reactions without purification.

EXAMPLE 66

Dansyl peptide 1,1,1-Trifluoro-3-(alanylalanylprolylamino)-4-methylpentan-2-one

To a suspension of 1,1,1-Trifluoro-3-(alanylalanylprolylamino)-4-methylpentan-2-one (50 mg, 0.11 mmol) in methylene chloride (1 ml), was added N-methylmorpholine (50 mg, 0.5 mmol). The solution was stirred for 5 min and dansyl chloride (50 mg) then added. The reaction mixture was stirred for 2 h at room temperature with the exclusion of light and then loaded directly onto a silica gel column (230-400 mesh) for purification. Elution with ethylacetate gave the dansylated peptide (48 mg, 0.07 mmol) in 68% yield.

EXAMPLE 67

N¹-(2-Methoxyethoxymethoxy-3,3-difluoro-1-isobutyl-5-hexenyl)-N²-isovaleryl valinamide

To 0.211 g of sodium hydride (55%, 4.83 mmol) in 3 ml of DMF at 0°C was added 1.8 g (4.6 mmol) of N¹-(2-hydroxy-3,3-difluoro-1-isobutyl-5-hexenyl)-N²-isovaleryl valinamide, in 5 ml of DMF. After stirring 0°C for 10 min., methoxyethoxymethylchloride (0.659 g, 5.29 mmol in 3 ml DMF) was added, the mixture stirred for 10 min. at 0°C and overnight at room temperature. Workup with water/Et₂O gave, after purification by flash chromatography (CHCl₃/Et₂O, 2:1) 1.4 g of the desired product.

EXAMPLE 68

N¹-(2-Methoxyethoxymethoxy-3,3-difluoro-1-isobutyl-4-carboxybutyl)-N²-isovaleryl valinamide

5 The above-named compound was prepared from N¹-2-methoxyethoxymethoxy-3,3-difluoro-1-isobutyl-5-hexenyl)-N²-isovaleryl valinamide by the procedure described in example 8 using equivalent proportions and conditions.

EXAMPLE 69

10 N¹-(2-Methoxyethoxymethoxy-3,3-difluoro-1-isobutyl-4-[1-(Isoamylaminocarbonyl)ethylmethylamino carbonylbutyl]-N²-isovaleryl valinamide

15 The above-named compound was prepared from N¹-(2-methoxyethoxymethoxy-3,3-difluoro-1-isobutyl-4-carboxybutyl)-N²-isovaleryl valinamide by the procedure described in example 49.

Proportions: 1.50 g (3.02 mmol) peptide acid in 10 ml THF, 0.306 g (0.33 ml, 3.02 mmol) N-methyl morpholine, 0.412 g (3.02 mmol) isobutylchloroformate, N-methyl alanine isamylamide hydrochloride (0.63 g, 3.02 mmol) and N-methyl morpholine (0.306 g, 3.02 mmol) in 5 ml THF. Flash chromatography (EtOAc/Pentane, 2:1) gives 0.3 g of the above-named compound. From 1.50 g of the peptide acid, using the procedure of example 49 there is produced, after purification by flash chromatography, 0.39 g of the desired product.

20
25

EXAMPLE 70

N¹-(2-Hydroxy-3,3-difluoro-1-isobutyl-4[[1-(isoamylamino
carbonyl)ethyl]-methylamino]carbonylbutyl)-N²-isovaleryl
valinamide

5 A mixture of 0.3 g (0.46 mmol) of N¹-(2-
methoxyethoxymethoxy-3,3-difluoro-1-isobutyl-4-[[1-
(isoamylaminocarbonyl)ethyl]-methylamino]carbonylbutyl)-N²-
isovaleryl valinamide and 0.52 g (2.31 mmol) of ZnBr₂ in 3
ml CH₂Cl₂ was stirred for 24 h at room temperature. Flash
10 chromatography (EtOAc) gives 0.11 g of the above-named
alcohol.

EXAMPLE 71

N¹-(2-Oxo-3,3-difluoro-1-isobutyl-4[[1-
(isoamylamino)ethyl]-methylamino]carbonylbutyl)-N²-
15 isovaleryl valinamide

The above-named compound was prepared from N¹-(2-
Hydroxy-3,3-difluoro-1-isobutyl-4[[1-
(isoamylamino)ethyl]-methylamino]carbonylbutyl)-N²-
isovaleryl valinamide by the procedure described in example
20 7.

Proportions: Oxalylchloride (0.176 mmol, 0.0224 mg) in 0.5
ml CH₂Cl₂, 0.0275 mg (0.352 mmol) DMSO, 90 mg of alcohol in
1.5 ml CH₂Cl₂ (at -55°C) 0.081 g Et₃N (0.8 mmol).

Flash chromatography (EtOAc) gave 0.02 g of above-
25 named compound.

EXAMPLE 72

4-N-tert-butoxycarbonylamino 2,2-difluoro 3-hydroxy 5-methyl hexanoic acid, ethyl ester

5 The title compound was prepared in 35% yield from L-BOC valinal using the same procedure as for the ester, described in example 32. $R_f = 0.52$ (EtOAc/C₆H₁₂ 1:1)

EXAMPLE 73

4-Amino 2,2-difluoro 3-hydroxy 5-methyl hexanoic acid, ethyl ester hydrochloride

10 The Boc protecting group of the alcohol of example 72 is cleaved using the same procedure as for the amide described in example 35, mp 182°C.

EXAMPLE 74

15 4-[methoxysuccinyl L-alanyl-L-alanyl-L-prolyllamino 2,2-difluoro 3-hydroxy 5-methyl hexenoic acid, ethyl ester

To a stirred solution of 0.371 g (1 mmol) of MeOSuc-L-Ala-L-Ala-L-ProOH in dry acetonitrile (10 ml) under nitrogen was added 0.106 g (1.05 mmol) of N-methylmorpholine. The resultant solution was cooled to -20°C.

20 Isobutyl chloroformate (0.136g, 1 mmol) was added to the cooled reaction mixture. After 10 min. a solution of 0.275 g (1.05 mmol) of 4-amino 2,2-difluoro 3-hydroxy 5-methyl hexanoic acid, ethyl ester hydrochloride and 0.106 g (1.05 mmol) of N-methylmorpholine in dry DMF (2 ml) was added to

25 the cooled mixture. The reaction mixture was stirred at -20°C for 4 hours and then allowed to warm to room temperature. After stirring 15 hours at room temperature the mixture is concentrated and placed under high vacuum at 40°C to remove all the DMF. Chromatography (silica gel,

0195212

ethyl acetate/acetone 7:3) yielded the expected alcohol in 85% yield. Rf: 0.38 (ethyl acetate/acetone 1:1).

EXAMPLE 75

5 4-[Methoxysuccinyl-L-alanyl-L-alanyl-L-prolyllamino 2,2-difluoro 5-methyl 3-oxo hexanoic acid, ethyl ester

The title compound was obtained in 65% yield from the alcohol of example 74 using the procedure described in example 37, mp: 96-97°C.

The foregoing describes in detail the generic and specific aspects of the scope of the invention as well as the manner of making and using the invention. In addition thereto, although such procedures are known in the art, references setting forth state of art procedures by which the compounds may be evaluated for their biochemical effects is also included herein.

For example, human elastase is assayed in vitro using chromophoric peptides, succinylalanylalanylalanyl-p-nitro-anilide (A1), methoxysuccinylalanylalanylprolylvalyl-p-nitroanilide (A2), and others, all of which are available commercially. The assay buffer, pH 8.0, and assay techniques are similar to those described by Lottenberg et al. (A3, A4). Enzyme is purified from human sputum (A5), although recently it has become commercially available. Kinetic characterization of immediate inhibitors is by means of the Dixon plot (A6), whereas the characterization of slow- and/or tight-binding inhibitors used data analysis techniques reviewed by Williams and Morrison (A7).

Similarly, the other proteases are assayed and effects of inhibitors are assessed in vitro by similar spectroscopic techniques: cathepsin G (A2); thrombin (A3); chymotrypsin (A8); trypsin (A9); plasmin (A3); C1 esterase (A10); urokinase (A3); plasminogen activator (A11); acrosin (A12); beta-lactamase (A13); cathepsin B (A14); pepsin (A15); cathepsin D (A16) and leucine aminopeptidase (A17). Pseudomonas elastase is measured in a coupled assay procedure using a human elastase substrate and microsomal aminopeptidase.

Radiometric assays of angiotensin I-converting enzyme and enkephalinase and their inhibitors are based on the procedure of Ryan (A18) and use tritiated substrate pur-

chased from Ventrex Laboratories, Inc. Radioimmunoassay is used for studies with renin (A19). C3-convertase is measured as described by Tack et al. (A20).

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By following the techniques referenced above, as well as by utilization of other known techniques, as well as by comparison with compounds known to be useful for treatment of the above-mentioned disease states, it is believed that adequate material is available to enable one of ordinary skill in the art to practice the invention. Of course, in the end-use application of the compounds of this invention, the compounds are preferably formulated into suitable pharmaceutical preparations such as tablets, capsules or elixers, for oral administration or in sterile solutions or suspensions for parenteral administration. The compounds of this invention can be administered to patients (animals and human) in need of such treatment in a dosage range of 0.01-10 mg per kg of body weight per day. As stated above, the dose will vary depending on severity of disease, weight of patient and other factors which a person skilled in the art will recognize.

Typically the compounds described above are formulated into pharmaceutical compositions as discussed below.

About 10 to 500 mg of a compound or mixture of compounds of Formula I or a physiologically acceptable salt is compounded with a physiologically acceptable vehicle, carrier, excipient, binder, preservative, stabilizer, flavor, etc., in a unit dosage form as called for by accepted pharmaceutical practice. The amount of active substance in these compositions or preparations is such that a suitable dosage in the range indicated is obtained.

Illustrative of the adjuvants which may be incorporated in tablets, capsules and the like are the following: a binder such as gum tragacanth, acacia, corn starch or gelatin; an excipient such as microcrystalline cellulose; a
5 disintegrating agent such as corn starch, pregelatinized starch, alginic acid and the like; a lubricant such as magnesium stearate; a sweetening agent such as sucrose, lactose or saccharin; a flavoring agent such as peppermint, oil of wintergreen or cherry. When the dosage unit form is
10 a capsule, it may contain in addition to materials of the above type, a liquid carrier such as fatty oil. Various other materials may be present as coatings or to otherwise modify the physical form of the dosage unit. For instance, tablets may be coated with shellac, sugar or both. A syrup
15 or elixer may contain the active compound, sucrose as a sweetening agent, methyl and propyl parabens as preservatives, a dye and a flavoring such as cherry or orange flavor.

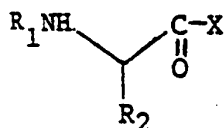
Sterile compositions for injection can be formulated
20 according to conventional pharmaceutical practice by dissolving or suspending the active substance in a vehicle such as water for injection, a naturally occurring vegetable oil like sesame oil, coconut oil, peanut oil, cottonseed oil, etc. or a synthetic fatty vehicle like
25 ethyl oleate or the like. Buffers, preservatives, antioxidants and the like can be incorporated as required.

While the invention has been described in connection with specific embodiments thereof, it will be understood that it is capable of further modifications and this
30 application is intended to cover any variations, uses, or adaptations of the invention following, in general, the principles of the invention and including such departures

from the present disclosure as come within known or customary practice within the art to which the invention pertains and as may be applied to the essential features hereinbefore set forth, and as follows in the scope of the
5 appended claims.

Claims:

1. An activated electrophilic ketone-bearing
peptidase inhibitor of the formula



the hydrates thereof, and the pharmaceutically acceptable salts thereof wherein

R_1 may be hydrogen, an amino protecting group selected from Group K, an α -amino acid or a peptide comprised of a number of α -amino acid building blocks, each of said α -amino acid or peptide optionally bearing an amino protecting group selected from Group K,

R_2 is the R group side chain of an α -amino acid building block,

X is X_1 or X_2 wherein

X_1 is $-\text{CF}_3$, $-\text{CF}_2\text{H}$, CO_2R_3 or $-\text{CONHR}_3$,

X_2 is $-\text{CF}_2\text{CHCR}_5\text{Y}$, $-\text{CF}_2\text{CR}_5\text{Y}$ or $-\text{CR}_5\text{Y}$,

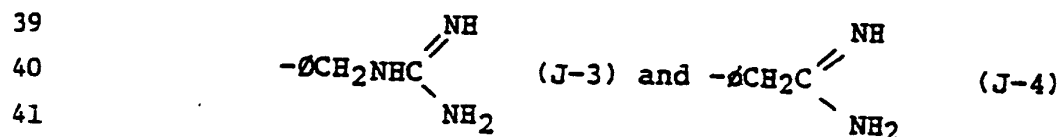
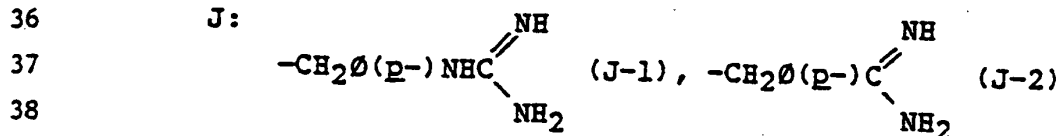


R_3 is hydrogen, C_1 -4 straight or branched alkyl, phenyl, benzyl, cyclohexyl or cyclohexylmethyl,

R_4 is the R group side chain of an α -amino acid building block,

23 R₅ is an α-amino acid or peptide building block, or deleted
 24 Y is -NHR₃ or -OR₃, said α-amino building blocks being
 25 selected from Groups A, B, C, D, E, F, G, K and J,
 26 members of these Groups being

27 Group A: Lys and Arg,
 28 B: Glu and Asp,
 29 C: Ser, Thr, Gln, Asn, Cys and His,
 30 D: Pro, Ind
 31 E: Ala, Leu, Ile, Val, n-Val, Met and n-Leu, and
 32 their N-methyl derivatives,
 33 F: Phe, Tyr and Trp, Nal(1) and their N-methyl
 34 derivatives,
 35 G: Gly, Sar
 36 J:



42 with ϕ representing phenyl

43 K: Acetyl (Ac), Succinyl (Suc), Benzoyl (Bz), t-
 44 Butyloxycarbonyl (Boc), Carbobenzoyl (CBz), Tosyl
 45 (Ts), Dansyl (DNS), Isovaleryl (Iva),
 46 Methoxysuccinyl (MeOSuc), 1-Adamantanesulphonyl
 47 (AdSO₂), 1-Adamantaneacetyl (AcAc), 2-
 48 Carboxybenzoyl (2-CB) and such other terminal
 49 amino protecting groups which are functionally
 50 equivalent thereto.

1 2. A compound of claim 1 useful for inhibiting human
 2 leukocyte elastase wherein

3 R₁ is -P₂P₃P₄P₅ wherein P₂ is selected from Groups D, E or
 4 F,
 5 P₃ is selected from Groups D or E,
 6 P₄ is selected from Group E or is zero,
 7 R₅ is selected from Group K,
 8 R₂ is the R group side chain of an α-amino acid of Groups E
 9 and G,
 10 X is either X₁ or X₂ wherein
 11 R₄ is the R group side chain of an α-amino acid of Groups E
 12 and G,
 13 R₅ is a building block having its α-amino acids selected
 14 from Groups E and G, and
 15 Y is NH₂.

1 3. A compound of claim 2 having the formula
 2 MeOSuc-Ala-Ile-Pro-Val[CF₂-Ala]AlaNH₂.

1 4. A compound of claim 1 useful for inhibiting
 2 Cathepsin G wherein X₁, X₂, R₃, R₄, R₅ and Y are as defined
 3 in claim 2,
 4 R₁ is P₂P₃P₄P₅ wherein
 5 P₂ is selected from Groups D, E, G or K,
 6 P₃ is selected from Groups E or G or is deleted,
 7 P₄ is selected from Groups E or G or is deleted,
 8 P₅ is selected from Group K,
 9 R₂ is the R group side chain of Groups E and F.

1 5. A compound of claim 4 having the formula
 2 Suc-Ala-Ala-Pro-Phe-CF₃.

1 6. A compound of claim 1 useful for inhibiting
 2 thrombin with X₁, X₂, R₃ being as defined in claim 1,
 3 R₂ is an R group side chain of Groups A and J,
 4 R₄ is an R group side chain of Group C or G,
 5 R₅ has its α-amino acids selected from Groups E, D or is
 6 zero,

- 7 R₁ is (a) -P₂P₃, (b) -P₂ or (c) -P₂P₃P₄ wherein
8 (a) P₂ is selected from Groups E and F, P₃ is
9 selected from Group F, each P₃ in its D
10 configuration,
11 (b) P₂ is selected from Group K,
12 (c) P₂ is selected from Group E, P₃ is selected from
13 Groups G or E, and P₄ is selected from Groups G
14 or E or is zero.

1 7. A compound of claim 6 having the formula
2 D-Phe-Pro-Arg-CF₃.

1 8. A compound of claim 1 useful for inhibiting
2 chymotrypsin wherein X₁, X₂, R₃, R₄, R₅ and Y are as
3 defined in claim 2,
4 R₁ is -P₂P₃P₄P₅,
5 P₂ is selected from Groups D, E, G or H,
6 P₃ is selected from Groups E or G or is zero,
7 P₄ is selected from Groups E or G or is zero,
8 P₅ is selected from Group K or is zero when P₂ is H.
9 R₂ is an R group side chain of Groups E or F.

1 9. A compound of claim 8 selected from the group
2 Bz-Phe-CF₃, BzPhe-COOH, BzPheCOOMe, BzTyrCF₃, Bz-Tyr-COOMe.

1 10. A compound of claim 1 useful for inhibiting
2 trypsin wherein X₁, X₂, R₃, Y, R₂, R₄, R₅ and R₁ are as
3 defined in claim 1.

1 11. A compound of claim 10 having the formula
2 Bz-Arg-COOH, Bz-Arg-CO₂CH₃, Bz-Arg-CF₃.

1 12. A compound of claim 1 useful for inhibiting
2 plasmin wherein
3 X is X₁,

4 R₁ is -P₂P₃P₄ wherein

5 P₂ is selected from Group F,

6 P₃ is selected from Groups B or F, and

7 P₄ is selected from Group K.

8 R₂ is the R group side chain of Groups A or J.

1 13. A compound of claim 12 selected from the group
2 DNS-Glu-Phe-Lys-COOH, DNS-Glu-Phe-Lys-CF₃, or
3 DNS-Glu-Phe-Lys-COOMe.

1 14. A compound of claim 1 useful for inhibiting C₁-
2 esterase with X₁, X₂, R₅ and R₃ as defined in claim 1,
3 Y is NH₂,
4 R₁ is -P₂P₃ wherein
5 P₂ is selected from Groups E, G, D, C, F, A or B, and
6 P₃ is selected from Group K,
7 R₂ is the R group side chain of Groups A and J,
8 R₄ is the R group side chain of Group E.

1 15. A compound of claim 14 selected from the group
2 consisting of CBZ-Ala-Arg-CF₃, CBZ-Ala-Arg-COOH and
3 CBZ-Ala-Arg-COOMe.

1 16. A compound of claim 1 useful for inhibiting C₃-
2 convertase with X₁, X₂ and R₃ being as defined in claim 1,
3 Y is OR₃,
4 R₁ is -P₂P₃P₄ with
5 P₂ being selected from Groups E or F,
6 P₃ is selected from Groups E or F,
7 P₄ is selected from Group K,
8 R₂ is the R group side chain selected from Groups A or J,
9 R₄ is selected from Group E, and
10 R₅ is zero.

1 17. A compound of claim 16 selected from the group
2 consisting of Bz-Leu-Ala-Arg-CF₂COOCH₂Ø,
3 Bz-Leu-Ala-Arg-CF₂-Ala-OCH₂Ø, Bz-Leu-Ala-Arg-COOCH₂Ø.

1 18. A compound of claim 1 useful for inhibiting
2 Urokinase wherein X₁ and X₂ are as defined in claim 1,
3 Y is NH₂,
4 R₁ is -P₂P₃ wherein
5 P₂ is selected for Groups E and G,
6 P₃ is selected from Group B,
7 R₂ is the R group side chain selected from Groups A and J,
8 and each of R₄ and R₅ is selected from Group E.

1 19. A compound of claim 18 selected from the group
2 consisting of Glu-Gly-Arg-CF₂H, Glu-Gly-Arg-CF₃,
3 Glu-Gly-Arg-COOH and Glu-Gly-Arg-CONH₂.

1 20. A compound of claim 1 useful for inhibiting
2 plasminogen activator with X₁ and X₂ being as defined in
3 claim 1,
4 Y is NH₂,
5 R₁ is -P₂P₃P₄ wherein
6 P₂ is Gly,
7 P₃ is selected from Group B, and
8 P₄ is selected from Group K,
9 R₂ is the R group side chain selected from Groups A and J,
10 R₄ is the R group side chain selected from Group E,
11 R₅ is selected from Group E.

1 21. A compound of claim 20 selected from the group
2 consisting of DNS-Glu-Gly-Arg-COOMe, DNS-Glu-Gly-Arg-CF₃,
3 DNS-Glu-Gly-Arg-COOH.

1 22. A compound of claim 1 useful for inhibiting
2 acrosin wherein X₁ and X₂ are as defined in claim 1,
3 Y is NH₂,

1 R₄ is an R group side chain selected from Group E,
 2 R₅ is selected from Group E,
 3 R₁ is -P₂P₃P₄ wherein
 4 P₂ is selected from Group E,
 5 P₃ is selected from Group E, and
 6 P₄ is selected from Group K,
 7 R₂ is an R group side chain selected from Groups A and J.

1 23. A compound of claim 22 selected from the group
 2 consisting of Boc-Leu-Leu-Arg-CF₂H, Boc-Leu-Leu-Arg-CF₃ and
 3 Boc-Leu-Leu-Arg-COOH.

1 24. A compound of claim 1 useful for inhibiting B-
 2 lactamase wherein X is X₁, as defined in claim 1 with the
 3 proviso that the P₁ carbonyl moiety may also exist in its
 4 reduced form,

5
 6 R₁ is P₂ which is selected from Group K or is $\text{OCH}_2\overset{\text{O}}{\parallel}\text{C}$.
 7 R₂ is an R group side chain selected from Groups E, G and
 8 C.

1 25. A compound of claim 24 selected from the group
 2 consisting of $\text{OCH}_2\text{CONHCH}_2\text{COCF}_3$, $\text{OCH}_2\text{CONH}_2\text{CHOHCF}_3$,
 3 $\text{OCH}_2\text{CONHCH}_2\text{COCOOH}$, $\text{OCH}_2\text{CONHCH}_2\text{COCOOME}$, $\text{OCH}_2\text{CONHCH}_2\text{CHOHCOOH}$
 4 and $\text{OCH}_2\text{CONHCH}_2\text{CHOHCOOME}$.

1 26. A compound of claim 1 useful for inhibiting D-
 2 Ala-D-Ala carboxypeptidase wherein X₁, X₂ and R₃ are as
 3 defined in claim 1,
 4 Y is OH or OR₃
 5 R₅ is deleted,
 6 R₄ is D-Ala,
 7 R₂ is the R group side chain of D-Ala and
 8 R₁ is -P₂P₃ wherein
 9 P₂ is selected from Groups E, C and (N α , ϵ)-di-AcLys, and
 10 P₃ is selected from Group K.

28. A compound of claim 1 useful for inhibiting cathepsin B wherein X_1 , X_2 , R_3 are as defined in claim 1, Y is OH,

R_5 is selected from Groups E, F or G,

R_4 is the R group side chain selected from Group E,

R_1 is (a) $-P_2P_3$ or (b) $-P_2P_3P_4$ wherein

(a) P_2 is selected from Groups E and F,

P_3 is selected from Group K,

(b) P_2 is selected from Groups E and F,

P_3 is selected from Groups E and F, and

P_4 is selected from Group K,

R_2 is the R group side chain selected from Groups A, J or Thr-CH₂O.

29. A compound of claim 28 selected from the group consisting of Ac-Leu-Leu-Arg†CF₂-Leu†Gly-OH, CBZ-Phe-Arg†CF₂-Leu†Gly-OH and CBZ-Phe-Thr†CF₂-Leu†Gly-OH.

30. A compound of claim 1 useful as inhibitor of renin wherein X_1 , X_2 and R_3 are as defined in claim 1, Y is OH wherein R_4 is the R group side chain selected from Groups E, F or G, R_5 is $P_2^1 P_3^1 P_4^1$ wherein P_2^1 is selected from Groups E, F or is deleted, P_3^1 is selected from Groups E, F or is deleted, P_4^1 is selected from Groups E, C, F or is deleted,

10 R₂ is the R group side chain selected from Groups E or F,
 11 or is cyclohexylmethyl,
 12 R₁ is -P₂P₃P₄P₅P₆ wherein
 13 P₂ is selected from Groups E, C or F,
 14 P₃ is selected from Groups E or F,
 15 P₄ is selected from Groups E, D or F or is deleted,
 16 P₅ is selected from Groups E, C or F or is deleted,
 17 and
 18 P₆ is selected from Group K.

1 31. A compound of claim 30 selected from the group
 2 consisting of MeOSuc-His-Pro-Phe-His-CF₂-Val-Ile-His-OH,
 3 MeOSuc-Pro-Phe-His-CF₂-Val-Ile-His-OH,
 4 MeOSuc-His-Phe-His-CF₂-Val-Ile-His-OH,
 5 MeOSuc-His-Pro-Phe-His-CF₂-Val-Ile-OH,
 6 MeOSuc-His-Pro-Phe-His-CF₂-Val-His-OH.

1 32. A compound of claim 1 useful for inhibiting
 2 pepsin wherein X₁, X₂ and R₃ being as defined in claim 1
 3 wherein
 4 R₄ is an R group side chain selected from Groups E, G or F,
 5 R₅ is selected from Groups E or F and
 6 Y is NHCH₂CH₂CH(CH₃)₂ or -NHCH₂CH(CH₃)₂,
 7 R₂ is an R group side chain selected from Groups E or F,
 8 R₁ is -P₂P₃P₄ wherein
 9 P₂ is selected from Groups E or F,
 10 P₃ is selected from Groups E or F,
 11 P₄ is selected from Group K.

1 33. A compound of claim 32 selected from the group
 2 consisting of Iva-Val-Val-Leu-CF₂-C(=O)NHCH(CH₃)₂ and
 3 Iva-Val-Val-Leu-CH₂-Gly-AlaNHCH(CH₃)₂.
 4

1 34. A compound of claim 1 useful for inhibiting cath-
2 epsin D wherein X_1 , X_2 and R_3 are as defined in claim 1,
3 Y is $-\text{NH}(\text{CH}_2)_2\text{CH}(\text{CH}_3)_2$ or $-\text{NHCH}_2\text{CH}(\text{CH}_3)_2$,
4 R_4 is an R group side chain selected from Groups E or F,
5 R_5 is selected from Groups E or F,
6 R_2 is an R group side chain selected from Groups E or F,
7 R_1 is $-\text{P}_2\text{P}_3\text{P}_4$ wherein
8 P_2 is selected from Groups E or F,
9 P_3 is selected from Groups E or F, and
10 P_4 is selected from Group K.

1 35. A compound of claim 34 selected from the group
2 consisting of CBZ-Val-Val-Phe- CF_2H , CBZ-Val-Val-Phe- CF_3 ,
3 CBZ-Val-Val-Phe- CF_2 -Phe-AlaNH $(\text{CH}_2)_2\text{CH}(\text{CH}_3)_2$,
4 CBZ-Val-Val-Phe- CF_2 -Phe-AlaNHCH $_2\text{CH}(\text{CH}_3)_2$.

1 36. A compound of claim 1 useful as an ACE inhibitor
2 wherein X is X_2 wherein
3 R_4 is an R group side chain selected from Groups E or G,
4 R_5 is selected from Groups A, B, C, D, E, F and G and Y is
5 OH,
6 R_2 is an R group side chain selected from Groups E, F or G,
7 and
8 R_1 is selected from Group K.

1 37. A compound of claim 36 having the formula
2 Bz-Phe[CF_2 -Gly]Pro-OH.

1 38. A compound of claim 1 useful for inhibiting
2 enkephalinase wherein X is X_2 wherein
3 R_4 is an R group side chain selected from Group E or F,
4 R_5 is selected from Groups E or F or is zero with the
5 proviso that when R_5 is zero, Y is $-\text{NH}_2$ and when it is
6 not zero, Y is $-\text{NH}_2$ or OH,
7 R_2 is Gly, and

8 R₁ is -P₂P₃ wherein
9 P₂ is Gly, and
10 P₃ is selected from Group F or is zero.

1 39. A compound of claim 38 selected from the group
2 consisting of Tyr-Gly-Gly-[CF₂-Phe]-Met-OH,
3 Tyr-Gly-Gly-[CF₂-Phe]-NH₂.

1 40. A compound of claim 1 useful as inhibitor of
2 pseudomonas elastase wherein X is X₂, wherein
3 R₄ is an R group side chain selected from Groups E or F,
4 R₅ is selected from Groups E or G and Y is NH₂,
5 R₂ is an R group side chain selected from Groups E and G,
6 and
7 R₁ is -P₂P₃, wherein
8 P₂ is selected from Group E and
9 P₃ is selected from Group K.

1 41. A compound of claim 40 having the formula
2 MeOSuc-Ala-Ala[CF₂-Ile]Ala-NH₂.

1 42. A compound of claim 1 useful for inhibiting
2 leucine aminopeptidase wherein X₁, X₂ and R₃ are as defined
3 in claim 1,
4 R₄ is an R group selected from any Group except H and
5 R₅ is selected from any Group except H,
6 Y is NH₂,
7 R₁ is hydrogen,
8 R₂ is an R group selected from Groups E or F.

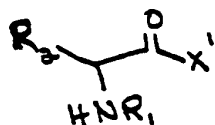
1 43. A compound of claim 42 selected from the group
2 consisting of Leu-CF₃, LeuCOOH, Leu[CF₂-Ala]AlaNH₂, and
3 LeuCOOMe.

1 44. A compound of claim 1 useful as an inhibitor of
2 kallikreins wherein X is X₁, R₂ is Arg, R₁ is a peptide
3 P₂P₃ with
4 P₂ being selected from Groups F and E,
5 P₃ being selected from Groups C, E or F.

1 45. A compound of claim 44 selected from the group
2 consisting of

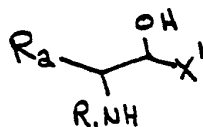
3 D-Pro-Phe-Arg-CF₂H,
4 D-Pro-Phe-Arg-CF₃,
5 D-Pro-Phe-Arg-CO₂H,
6 D-Pro-Phe-Arg-CONH₂

1 46. The process for producing compounds of the
2 formula



(A)

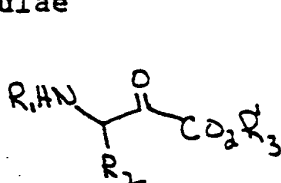
4 and the hydrate thereof
5 which comprises effecting a Swern oxidation on a
6 compound of the formula



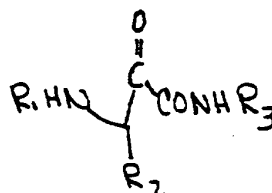
(B)

8 wherein X^1 is CF_2H or CF_3 , and R_1 and R_2 are as
9 defined in claim 1, and optionally removing any
10 protecting group.

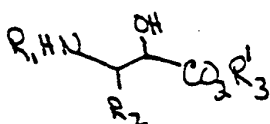
1 47. The process for preparing compounds of the
2 formulae



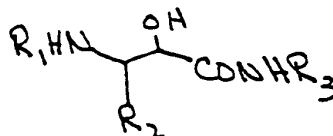
and

(C₁ and C₂)

4 and the hydrates thereof
5 which comprises effecting a Swern oxidation on
6 compounds of the formulae

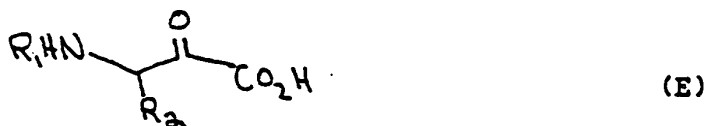


and

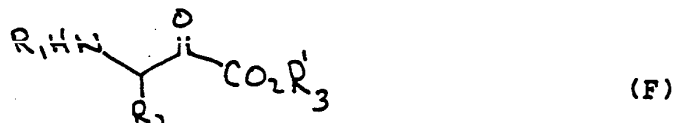
(D₁ and D₂)

wherein R_3 is as defined in claim 1, R_3' is R_3 except H, and R_1 and R_2 are as defined in claim 1, and optionally removing any protecting group.

48. A process for preparing a compound of the formula

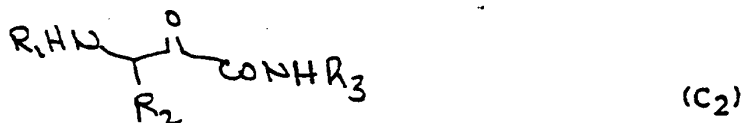


which comprises treating a compound of the formula

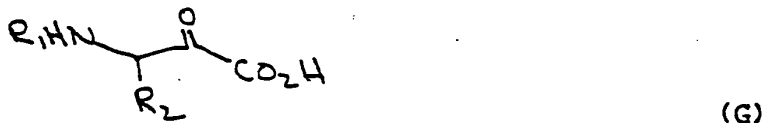


with a base, wherein R_1 , R_2 and R_3' are as defined in claim 47.

49. A process for preparing a compound of the formula

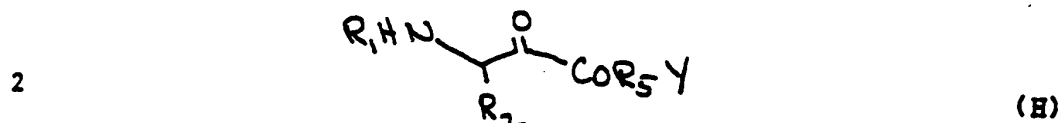


which comprises coupling $R_3\text{NH}_2$ to a compound of formula

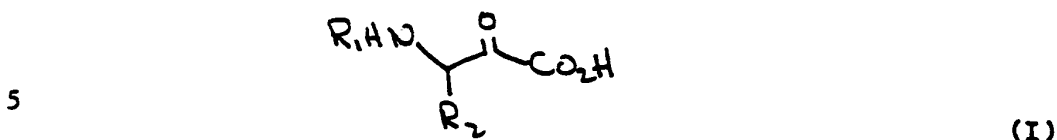


according to standard coupling procedures wherein R_1 , R_2 and R_3 are as defined in claim 47, and optionally removing any protecting group.

1 50. A process for preparing a compound of the formula

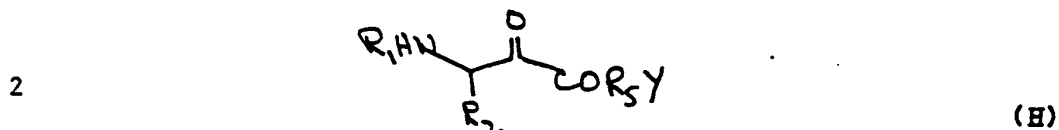


3 which comprises coupling R_5Y to a compound of the
4 formula

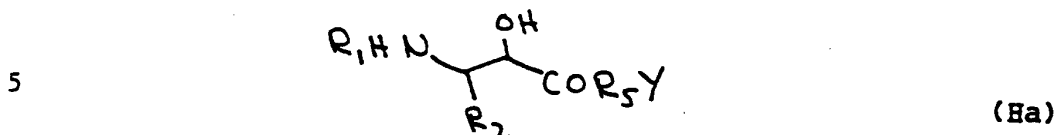


6 according to standard peptide chemistry procedures,
7 and optionally removing any protecting groups, wherein R_1 ,
8 R_2 , R_5Y are as defined in claim 1.

1 51. A process for preparing a compound of the formula

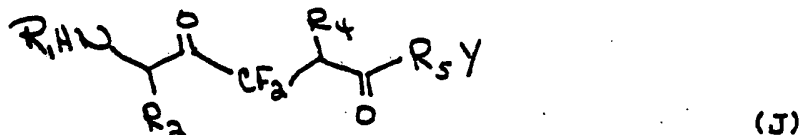


3 which comprises effecting a Swern oxidation on a
4 compound of the formula

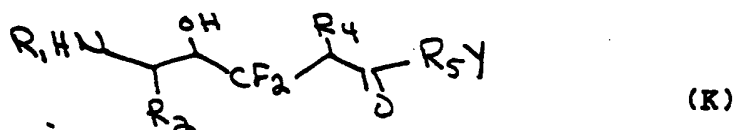


6 and optionally removing any protecting group, wherein
7 R_1 , R_2 and R_5Y are as defined in claim 1.

52. The process for preparing a compound of the formula

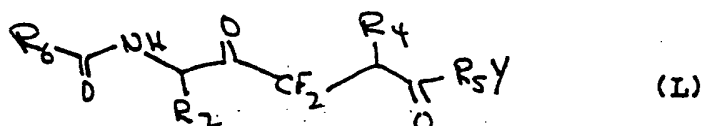


which comprises effecting a Swern oxidation on a compound of the formula

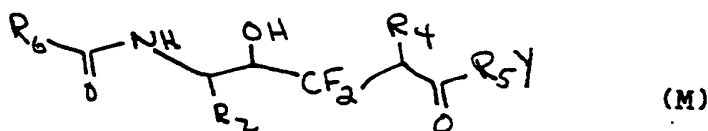


and optionally removing any protecting groups.

53. A process for preparing a compound of the formula

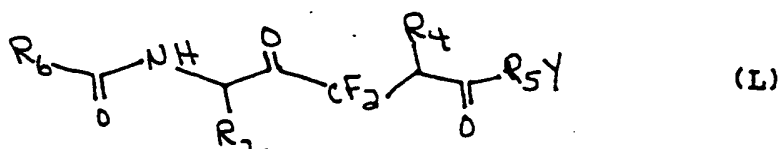


which comprises effecting a Swern oxidation on a compound of the formula

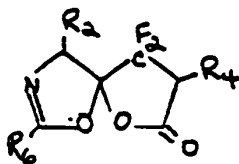


and optionally removing any protecting group wherein R_2 , R_4 , R_5Y are as defined in claim 1, and R_6 is phenyl, benzyl or other equivalently functioning group.

54. A process for preparing a compound of the formula



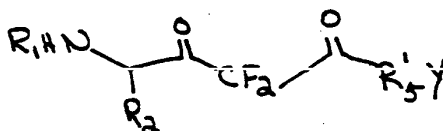
which comprises coupling R₅Y with a compound of the formula



(La)

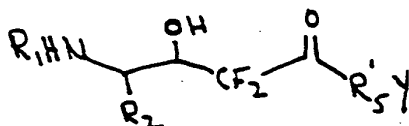
and optionally removing any protecting group, wherein R₂, R₄, R₅Y and R₆ are as defined in claim 53.

55. A process for preparing a compound of the formula



(M)

which comprises oxidizing a compound of the formula



(Ma)

according to Swern oxidation conditions or with pyridinium dichromate, and optionally removing any protecting group wherein R₁ and R₂ are as defined in claim 1 and R'₅Y is as defined for R₅Y except that R₅ optionally contains a protecting group.